

CURRENT DEBATES ON HEALTH SCIENCES

4

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Current Debates on Health Sciences 4

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Evaluation of Dose Distribution Verification of Treatment Plans by Using Electronic Portal Imaging Device at Different Gantry Angles in Intensity Modulated Radiation Therapy

Alper ÖZSEVEN¹

Introduction

Intensity Modulated Radiation Therapy (IMRT) is a radiation treatment that has become a standard therapy modality, allowing the target volume to be treated at desired doses compared to 3-dimensional conformal radiation therapy, while the normal tissues in the vicinity of the tumor are exposed to less radiation (Van Esch et al., 2004:223). Electronic portal imaging device (EPID) is a retractable component of linear accelerator used in radiation therapy and is one of the most functional equipment in terms of verifying the accuracy of the dose distribution in IMRT treatments as well as validating the position accuracy before the daily treatment of the patients (van Elmpt et al., 2008:289-290). In addition to the fact that IMRT is a complex treatment, the accuracy of treatment depends on many parameters that makes it vital to ensure patient-based quality assurance (QA) (Bailey et al., 2012:83); (van Zijtvelde et al., 2006:168); (Sharma et al., 2010:238). One of the most important parameters in LINACs that affecting the treatment dose distribution accuracy is the correct movement of the multi-leaf collimators (MLC), which form the treatment field depending on the shape of the target tumor (Chang et al., 2004:2091); (Richart et al., 2012:263). The studies can be found in the literature that describe in detail how EPID is used in pretreatment verification of IMRT (van Elmpt et al., 2008:290-294); (Jornet et al., 2014:381-382).

Gamma index analysis method has been used as a standard technique for patient-based pretreatment dose verification for many years. Gamma index analysis evaluation is performed by comparing the dose distribution fluence obtained from EPID with the dose distribution fluence calculated and estimated before the irradiation by the treatment planning system for a certain dose difference (DD) at a certain distance to agreement points (DTA) (Low et al., 1998:657); (Spezi et al., 2006:225); (Hussein et al., 2013:370); (Chaikh et al., 2016:32). While it is accepted as standard that to perform gamma analysis with 3% dose difference at 3mm (3%/3mm), it is possible to make comparisons at lower values as an evaluation criterion. Area gamma index <1 value above 95% is considered to be within the acceptance limit. Moreover, the value of area gamma <1 represents the passing rate of the total points for gamma analysis.

In routine practice, the verification of the dose accuracy of the treatment plans of the patients who will receive treatment with the IMRT modality is carried out with the technique of irradiating the EPID with treatment head (gantry) at a fixed angle, which is 0°. This procedure is also accepted as the reference verification position, as it allows faster results. On the other hand, approximately 64% of clinical centers perform the verification of dose distribution with the technique named as non-fixed gantry angles, which are predefined in treatment planning process (Nelms and Simon, 2007:81). Theoretically, it is expected that the treatment devices have the same dose output and therefore the same dose distribution at all gantry angles, however the MLC and

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EPID systems, which are the more sensitive parts of the LINACs to the gravity, are likely to behave differently depending on the angle. This can cause slight variations in the dose distributions (Chin et al.,2004:102); (Monti et al.,2013:205); (Cyriac et al.,2014:2).

The aim of this study is to compare the effects of pretreatment patient-based quality assurance in IMRT patients on the dose distribution of two different techniques, which are the irradiation technique at a fixed gantry angle, 0°, and the irradiation technique using gantry angles, which are predefined in the treatment planning procedure.

METHOD

In this study, 30 randomly selected treatment fields of patients treated with IMRT were compared using two different pretreatment quality assurance techniques (Figure 1). Treatment plans were created using the Eclipse treatment planning system on Varian DHX linear accelerator, with dose calculation by an anisotropic analytical algorithm (AAA). 6 MV x-rays were used. First, the dose distributions on the EPID were recorded by irradiating at a fixed angle at 0°. After that, irradiation was repeated using the non-fixed gantry angles of the fields in the treatment plan and the dose distributions on the EPID were recorded. First of all, by using the Eclipse Portal Dosimetry software, area gamma index <1 values from the resultant dose distributions obtained from EPID for each measurement technique and the predicted dose distributions estimated by the software were compared using the 3%/3mm evaluation criteria (Figure 2). Wilcoxon signed-rank test was used to evaluate the statistical significance. The p-value less than 0.05 was considered statistically significant for all statistical analyses.



Figure 1 – Experiment set-up: pretreatment verification with fixed gantry angle (a), pretreatment verification with non-fixed gantry angle (b)

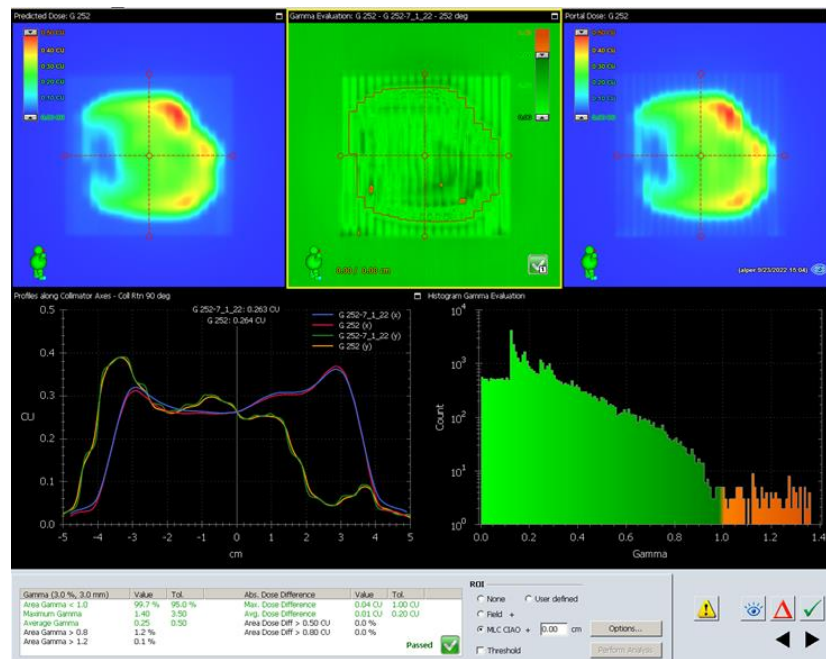


Figure 2 – Gamma analyses in the Eclipse Portal Dosimetry software: dose distributions obtained after irradiation (portal dose, upper left), estimation by the software (predicted dose, upper right), and blended view of comparison (upper middle)

In addition, by using Eclipse Portal Dosimetry software, fixed and non-fixed angle dose distributions, obtained as a result of irradiation were directly compared with each other using area gamma index <1 values in 3%/3mm and 1%/1mm evaluation criteria (Figure 3).

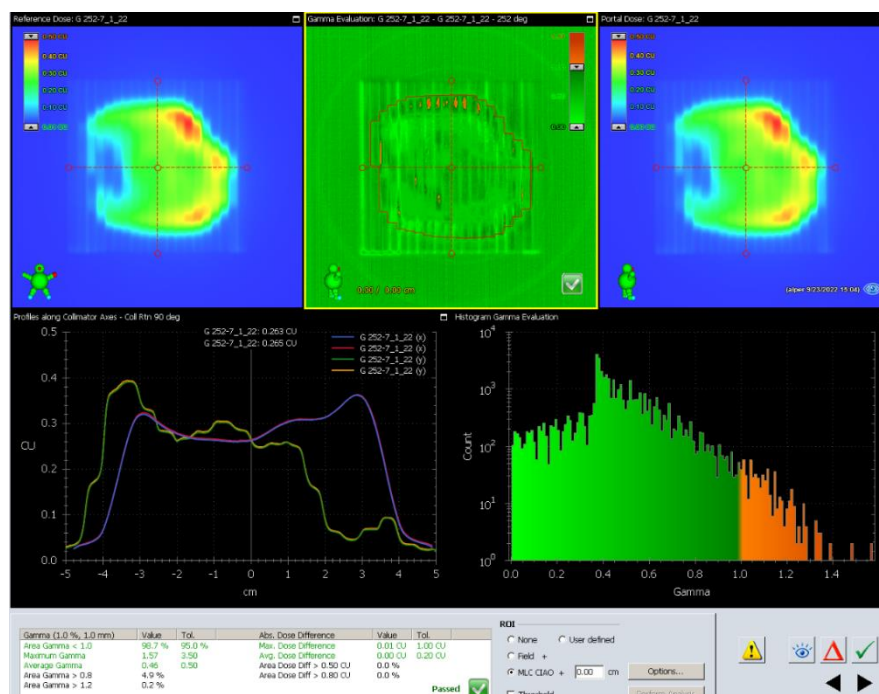


Figure 3 – Gamma analyses of direct comparison in Eclipse Portal Dosimetry software: fixed (upper left) and non-fixed (upper right) angled dose distributions obtained as a result of irradiation of EPID with 1%/1mm evaluation criteria and blended view of comparison (upper middle)

RESULTS and DISCUSSION

The area gamma index <1 median (min-max) values, which are obtained from EPID and the dose distributions estimated by the software with the 3%/3mm dose evaluation criterion for fixed and non-fixed gantry angles are 98.55% (93.70% - 100.00%) and 99.35% (93.90% - 100.00%), respectively ($p=0.03$) (Table 1).

In measurements where dose distributions were directly compared with each other, all the results were found to be 100% for 30 treatment fields in 3%/3mm evaluation criteria. Since it was a statistically constant value, the result was given as $100\% \pm 0\%$ with the notation of mean $\pm$ standard deviation. The area gamma index <1 median (min-max) values with the 1%/1 mm evaluation criteria are 98.75% (88% - 100.00%) (Table 1).

Table 1 – Area gamma <1 values as a result of patient-based quality assurance process with two different techniques

Gantry Angle Evaluation Criteria	Predicted Dose vs Portal Dose 0° Planned		Direct Comparison of the Fields (Planned vs Gantry 0°)	
	3%/3mm		1%/1mm	3%/3mm
Area Gamma <1 Value*	98.55% (93.70% - 100.00%)	99.35% (93.90% - 100.00%)	98.75% (88% - 100.00%)	$100\% \pm 0\%$
p value	0.03**			

** denotes that correlation is significant at the 0.05 level (2-tailed)

* median (min-max) or mean $\pm$ standard deviation which is applicable

In cases where dose differences in dose distributions need to be evaluated at a certain distance more precisely, this evaluation criterion may decrease to 2%/2mm or even 1%/1mm. In this study, the 3%/3mm criterion was chosen, because it is the reference evaluation criterion. Besides, to analyze the precision of the difference between fixed and non-fixed gantry angle techniques, the 1%/1mm criterion was chosen. The obtained results showed that there was no statistically significant difference for the 3%/3mm evaluation criterion in the comparison of fixed and non-fixed gantry angle techniques with the predicted dose distributions. In addition, the median values found for the two techniques were very close to each other and were higher than 95%, which is the lower limit for area gamma index <1 in clinical routine. On the other hand, no difference was found for the 3%/3mm criterion with the obtained results when these two techniques were directly compared with each other. In measurements where the evaluation criterion was 1%/1mm, the obtained median area gamma index <1 value was 98.75% and this value was higher than the lower limit of 95% and within the acceptance limit. It was remarkable that even with an evaluation criterion such as the 1%/1 mm, the attained result was within the acceptance limit.

In their multicenter study, Jornet et al. stated that pretreatment quality assurance irradiation of patients with head-neck and prostate cancer with fixed and non-fixed gantry angle, did not make a significant difference in the results and yielded values higher than 95%, which is the lower limit for area gamma index <1 (Jornet et al., 2014:386). Similarly, in the study of Cyriac et al., it was reported that the use of these two different techniques in the pretreatment quality assurance processes of patients with prostate cancer did not affect the outcomes. In addition to this, even if no dosimetric difference was found between these techniques; it was also noted that the pretreatment quality assurance process may vary from device to device (Cyriac et al., 2014:4). Monti et al. stated that the pretreatment quality assurance process at different angles caused a position change in the EPID system due to mechanical conditions. Nevertheless, no difference was measured on the dose distributions with this procedure (Monti et al., 2013:206).

CONCLUSIONS

As a consequence, although the difference was statistically significant for gamma analyzes with the predicted dose distributions, all the results were within the acceptance limits. In cases where the techniques were directly compared with each other, the results were within the acceptance limit, and the difference between them varied with regard to the selected evaluation criteria. With a significant time savings for each patient, the pretreatment IMRT QA may also be carried out at fixed (0°) gantry angles.

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Hyperhidrosis

Aykut ELİÇORA

Introduction

Sweating is a vital physiological process. Hyperhidrosis is defined as sweat secretion that affects the daily life of more people than physiological. It can be considered as a frequently recurring chronic autonomic disease. The most affected organs are the hands, feet and axillary region.

Hyperhidrosis can be primary or secondary. Primary hyperhidrosis is idiopathic and is usually bilaterally symmetrical. Secondary hyperhidrosis can be focal or generalized and is seen due to another underlying cause such as drug use, malignancies, alcohol, chronic pulmonary disease, congestive heart failure, endocrine metabolic disorders, febrile diseases. (Walling, 2011),(Hornberger & et al.,2004)

It has been shown that hyperhidrosis is inherited in an autosomal dominant manner. Positive family history was found in approximately 31.5-65% of them.

Thermoregulatory Sweating

Sweat glands are responsible for thermoregulation to maintain body temperature. It helps regulate body temperature in case of heat, fever or physical exertion. The sweating center is in the hypothalamus and is under the control of the frontal cortex. The sympathetic nervous system regulates thermoregulation by controlling the sweat glands and the circulation of the skin. Regulation is provided by the feedback loop of afferent somatosensory and central temperature sensitive neurons.(Schlereth & Dieterich & Birklein, 2009)

Emotional Sweating

Emotional sweating serves as a "feedback" signal from the emotional to the physical. This type of sweating is regulated by the neocortical and limbic centers. It also occurs with the stimulation of sweat glands in the hands, soles of feet and axilla, which are called "emotional sweating areas" in various emotional situations. Vasodilation accompanies stress-induced thermoregulatory sweating.(Bini & et al.,1980)

But it should be kept in mind that the distinction between thermoregulatory and emotional sweating is not an absolute distinction. It has been shown that they mutually affect each other

Surgical treatment.

Focal Hyperhidrosis

Focal hyperhidrosis is most commonly seen as axillary hyperhidrosis. Other forms seen include palmar and plantar hyperhidrosis. (Haider & Solish, 2005) focal hyperhidrosis first begins in puberty or adolescence. Although hyperhidrosis can continue for life, it may end spontaneously with advancing age. (Vlahovic, 2016) The sweating described here is at a level that cannot be explained by external factors. (Sammons & Khachemoun, 2017)

Three types of glands are involved in the realization of sweating. These are the eccrine, apocrine, and apoecrine sweat glands. Eccrine sweat glands begin to perform their duties from

the moment of birth. Glands that start to function after puberty are apocrine and apoeccrine glands. The increased sweating seen in primary hyperhidrosis is not caused by an increase in the number of these glands, but by an increase in their activity. (Vlahovic, 2016)

Excessive stimulation of the sympathetic nervous system is often cited as the cause of the increase in sweating, and therefore most treatments also show efficacy on the sympathetic nervous system. (Abu & Elhamami, 2021)

Excessive sweating, wet hands, staining on clothes, and odors that can be described as "unpleasant" by people may affect the social life of the person. In conclusion, if hyperhidrosis patients are examined outside of their physical distress, they suffer from higher rates of depressed mood; It is seen that they complain of reduced comfort and self-confidence in daily life conditions such as work, school, and leisure. (Sammons & Khachemoun, 2017)

Axillary Hyperhidrosis

Axillary hyperhidrosis, which is the most common form of primary hyperhidrosis, can be described as excessive sweating in the axilla. Although it manifests itself in puberty or adolescence, if treatment methods are not evaluated, it often continues for life. All three types of sweat glands are found in the axilla. As a result of the formation of conditions such as staining on clothes and odor that can be described as "unpleasant", problems may arise in the patient's relations with other people. (Sammons & Khachemoun, 2017)

Palmar Hyperhidrosis

Palmar hyperhidrosis, a form of focal hyperhidrosis, is a large amount of sweating on the hands that cannot be explained by external factors or emotional state, usually bilaterally symmetrical, and seen throughout the year. During this sweating, which is more severe in the palms, the color of the palm may be normal or close to red. Usually this sweating is cold and wet. (Romero, 2016) This sweating, which affects life socially, like axillary hyperhidrosis, can cause problems in the interpersonal relationships of the person and cause them to develop behaviors such as avoiding shaking hands. (Abu & Elhamami, 2021)

Plantar Hyperhidrosis

Focal hyperhidrosis that can occur on the soles of the feet is then called plantar hyperhidrosis. Affected patients talk about the anxiety felt while performing sports activities. As the reason for this, they show the necessity of changing foot accessories such as socks and shoes more than once during the day. (Vlahovic, 2016) As this sweat is metabolized by standing microfauna, malodorous sweating called bromohidrosis may occur. This condition, which is frequently seen in plantar hyperhidrosis, may cause the social complaints of patients to increase exponentially. (Rieger & Pedevilla & Lausecker, 2015) Plantar hyperhidrosis, which creates a moist environment on the feet, invites fungal infections, which are often referred to as foot, in addition to social complaints. The cutaneous infection that may occur may be of bacterial or viral origin apart from fungal. (Vlahovic, 2016)

Treatment

It can be divided into two as conservative and surgical options. However, full cure cannot be achieved. Treatment choices vary from patient to patient. As the main headings, treatment options can be listed as follows.

Topical treatments

Systemic treatments

Device-based treatments

Surgical treatments

HDSS (hyperhidrosis disease severity scale) score system is used in the selection of treatment. In HDSS, questions are asked to the patient to determine the severity of the disease and scoring is done. Scoring is used to determine treatment.

Topical Treatments

Antiperspirants

It is a frequently beneficial treatment method used in patients with HDSS step 1-2. Aluminum chloride hexahydrate is used in amounts ranging from 6.25-40%. Lower concentrations are suitable for sensitive skin, but efficacy also decreases with intensity.

This solution is used once a day. It is applied on dry skin, especially before going to bed at night. It destroys secretory cells and mechanically obstructs eccrine sweat glands. May cause temporary skin irritation. Aluminum zirconium tetrachlorohydrate (AZT), which has fewer side effects, is another treatment option.

Topical Anticholinergics

HDSS step 1-2 is a more commonly used treatment modality in craniofacial type hyperhidrosis. 1-2% glycopyrrolate is used as spray or solution once or twice a day. Typical anticholinergic effects such as xerostomia, mydriasis and urinary stasis can be seen as side effects. It has been shown to provide 96 percent success.

Cryotherapy

It has been observed that eccrine sweat glands are damaged by the cold application method using the nitrous oxide cryoprobe applicator.

Botulinum Toxin

Botulinum Toxin Type A is administered locally by intradermal injection. It provides temporary success.

Systemic Treatments

Anticholinergics

It is one of the most commonly used systemic treatments. Dose adjustment can be increased depending on the development of tolerance. It is not preferred by patients due to the excess of side effects.

It is contraindicated in cases of Myasthenia Gravis, Paralytic Ileus, pyloric stenosis. The most preferred oral anticholinergics are glycopyrrolate and oxybutynin. It is preferred in patients with HDSS score of 3 and above. Typical anticholinergic side effects (urinary stasis, constipation, dry mouth, tachycardia, etc.) are present.

Alpha adrenergic agonists

Clonidine, an alpha-2 adrenergic agonist, is used in the treatment of hyperhidrosis by reducing sympathetic outflow.

Calcium channel blockers and other oral medications

Helps reduce sweating by reducing the release of acetylcholine

Injectable treatments

Botulinum toxin is a neurotoxin that blocks acetylcholine. It is the most commonly used non-surgical treatment. It is especially preferred in the treatment of axillary hyperhidrosis. Botulinum Toxin A is divided into three. Onabotulinum, Abobotulinum, Incobotulinum.

Rimabotulinum is used in Botulinum Toxin B. In the side effects, hand muscle paralysis is seen especially in those who are injected into the hand.

Device Treatments

Microwave

Microwave thermolysis is used in the treatment of hyperhidrosis by causing local inflammation.

Iontophoresis

It is used in the treatment of palmoplantar hyperhidrosis. The exact mechanism has not been elucidated. An ionized substance is caused to damage the sweat glands through water and using electric current.

Radiofrequency

Microneedle radiofrequency is ablation of eccrine sweat glands by heating the hypodermis. Photodynamic therapy, Laser and Ultrasound are among the other treatment methods.

Surgical Treatments

Surgical methods are used in the failure of conventional treatment methods. Localized surgery can be performed in axillary hyperhidrosis. Endoscopic thoracic sympathectomy (thoracic 2nd and 3rd sympathetic ganglia are removed) is preferred in more severe cases. Even if the hyperhidrosis in the relevant region decreases, compensatory sweating in different regions (abdomen, back, hips) is common as a late complication. Other complications related to surgery are hemothorax pneumothorax horner syndrome ductus thoracic injury and N. phrenicus palsy, wound infection, skin necrosis, skin discoloration, and scar formation.

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Chest Wall Deformities

Aykut ELİÇORA

Introduction

Chest wall deformities are structural anomalies that affect the sternum, ribs and chest wall muscles and change the anatomical structure and/or function of the thorax. (Andreescu & et al., 2018) Chest wall deformities are a large group of anomalies that are seen with an incidence of 1/400 - 1/1000 in the general population and have a significant impact on the life of the patients. (Yüksel, 2015) Although the majority of the patients are in the childhood, it can also be encountered in the early adult period in the clinic. The incidence is higher in men. Although the etiology and/or pathogenesis cannot be clearly explained, family history is reported in most of the cases. (Yüksel, 2015)

The psychosocial and cosmetic concerns of the patients have made these deformities more important and remarkable in clinical practice, and have led to the development of different surgical and non-surgical treatment methods.

Pectus excavatum and pectus carinatum are the most common chest wall deformities. Apart from these, sternal cleft, ectopia cordis, Poland syndrome, Jeune syndrome are seen more rarely. Chest wall deformities may be accompanied by other musculoskeletal anomalies, cardiovascular, gastrointestinal and genitourinary anomalies.

Pectus Excavatum

Pectus excavatum or “funnelchest” (funnel chest), popularly known as “shoemaker's chest”, is the most common chest wall deformity. It accounts for approximately 90% of all chest wall deformities. It occurs in 1 in 400 live births and is more common in males (male/female ratio approximately 4:1). Although it is mostly noticed during birth or in the first years after birth, the severity of the deformity may increase with the growth of the child.

It is caused by the posterior depression of the sternum as a result of abnormal growth and inward collapse of the costal cartilages. It usually affects the middle and lower 1/3 of the sternum. Although the etiology is not clearly known, it has been reported that 37% of the cases have a family history. Previous studies have shown that inheritance in pectus excavatum is autosomal dominant; however, it is thought that autosomal recessive inheritance, X-linked inheritance and multifactorial inheritance may also play a role in genetic transmission. The coexistence of pectus excavatum with anomalies such as scoliosis, long extremities, high palate, mitral valve prolapse, cardiac arrhythmias shows that there may be a relationship between pectus excavatum and connective tissue disorders. (Andreescu & et al., 2018), (Creswick, et al., 2006) It can be seen in mild, moderate and severe forms in the clinic. Although the deformity can be centralized or asymmetrical, it is often asymmetrical, most commonly depression in the right hemithorax. It can rarely be seen in the axillary or lateral regions. The severity of symptoms may vary according to the degree of deformity. In mild cases, patients may be mostly asymptomatic. In severe cases, cardiac pressure, dyspnea, chest pain, palpitations, decreased exercise tolerance and syncope can be seen. In addition, patients with pectus excavatum often have conditions such as lack of self-confidence, decrease in social relations, anxiety and avoidance of social activities.

Abnormal changes can also be detected in the electrocardiography (ECG) and pulmonary function tests of patients with deformities. ST and T changes, P wave elevation, right axis deviation, right bundle branch block, left atrium and left ventricular hypertrophy can be seen

on ECG due to cardiac compression. Mitral valve prolapse can be detected by echocardiography in 18% of cases. Restrictive changes occur in pulmonary function tests. (Beiser, et al., 1972), (Cahill, Lees & Robertson, 1984) Compared to the general population, it was determined that there was a significant decrease in maximal voluntary ventilation in patients. Frequent recurrent pulmonary infections are also seen in approximately 30% of patients. After the treatment of the deformity, improvement in cardiac and pulmonary findings is observed. (Fonkalsrud, 2009) In addition, psychosocial recovery is also provided.

Pectus Carinatum

Pectus carinatum, popularly known as "pigeon chest", is the second most common chest wall deformity after pectus excavatum. It accounts for approximately 16% of chest wall deformities, more common in men than women (4:1). In contrast to pectus excavatum, it is rare for it to occur soon after birth, often not recognized until puberty. (Demirkaya & Cansever, 2018) As in pectus excavatum, the etiology is not known clearly, but a positive family history has been reported in 25-30% of the cases. Scoliosis, kyphosis, congenital heart diseases and various musculoskeletal anomalies may accompany.

Pectus carinatum is characterized by anterior protrusion of these structures as a result of abnormal growth of the costal cartilages and sternum. It can be classified into three types.

Chondrogladiolar type:

It is the anterior projection of the lower costal cartilages together with the sternum corpus. It is the most common type. It can be symmetrical (most common) or asymmetrical.

Mixed type

It is the type in which pectus excavatum and carinatum deformities are seen together.

Chondromanubrial type

It results from the anterior projection of the manubrium and the second and third rib cartilages and relative depression of the sternum corpus. It is rare.

Although functional disorders due to cardiac and pulmonary compression are less common than pectus excavatum, complaints such as chest pain, dyspnea with exertion, and an increase in respiratory tract infections can also be seen in patients with pectus carinatum. In addition, psychosocial and cosmetic concerns are the most important reasons for seeking treatment.

Treatment Methods In Pectus Deformities

The choice of the right treatment method should be carefully determined according to the shape and severity of the deformity, the age, gender and social status of the patient. Treatment protocols can be grouped under two headings as surgical and non-surgical methods. Traditional Ravitch method for pectus excavatum in surgical treatment protocols and Minimally Invasive Pectus Correction (MIRPE) operation "Nuss procedure" (Molik & et al., 2001); For pectus carinatum, traditional Ravitch method and today's minimally invasive equivalent "Abramson procedure" are applied. Vacuum bell for pectus excavatum and dynamic external compression (orthosis / corset treatment) for pectus carinatum are used in non-surgical protocols.

Surgical treatment

Surgical indications in the treatment of pectus excavatum can be evaluated in two groups as functional and psychological-cosmetic indications. The aim is to prevent the compression of the deformity on the heart and lungs, to allow the normal development of the thorax, to correct the posture and to improve the psychological state. The treatment protocol should be decided according to the shape of the deformity and the clinical condition of the patient, after a detailed

evaluation with the patient and his family. Although there is no clear consensus on the timing of surgery, the age range of 10-16 is considered to be the most appropriate. (Nuss & et al., 1998)

The Haller index (HI), which was presented as a result of Haller's studies in 1987, is the most commonly used index in the typing of pectus excavatum. It is the ratio of the transverse diameter at the deepest point of the deformity to the distance between the posterior surface of the sternum and the anterior aspect of the vertebrae at the same point in thorax computed tomography. In normal people, the HI is 2.5. An HI above 3.25 is an indication for surgery. (Haller, Kramer & Lietman, 1987)

The first surgical treatment for pectus deformities was performed by Ludwig Meyer in 1911 in a 16-year-old patient with pectus excavatum. Secondly, it was performed by Sauerbruch in 1913. Over the years, open surgical procedures with different techniques have continued to be applied and improvements have been made in the technique. The classical operation technique that will be used from the past to the present is the Ravitch Method. The first series of 8 cases, started by Mark M. Ravitch in 1947, was published in the April 1949 issue of the *Annals of Surgery* journal. (Ravitch, 1949) Complications after open surgery include atelectasis, pneumothorax, hemothorax, wound infections, pneumonia, intrathoracic organ injury, pericarditis, sternocostal joint dislocation, and recurrences in the late period.

Today, the most preferred method for pectus excavatum is the minimally invasive pectus excavatum correction operation (MIRPE), which was defined by Donald Nuss in 1988. In this technique, which is applied under general anesthesia, a nickel-steel alloy bar is placed videothoroscopically in the retrosternal area. The bar is removed at the end of 2-4 years. In the minimally invasive method, short operation time, minimal blood loss, earlier initiation of daily activities and shorter hospital stay are important advantages compared to other open surgical methods. Despite this, open surgical methods are still used in selected cases. The most serious complication of Nuss operation is cardiac injury. Early complications include pneumothorax, wound infection, hemothorax, nickel allergy, pleural effusion, pericarditis, and pneumonia, while late complications include bar and stabilizer dislocation, pain, infection, allergic reactions, hemothorax, recurrence, and overcorrection. With the developing surgical techniques and increasing surgical experience, the risk of complications and recurrence in pectus excavatum surgery has decreased.

The traditional method in the surgical treatment of pectus carinatum is the Ravitch method. It is an open technique that involves resection of deformed cartilages with or without sternal osteotomy, depending on the condition of the deformity. The Abramson Procedure, a modified version of the Nuss procedure for pectus carinatum, was described by Argentine thoracic surgeon Dr. Horacio Abramson in 2005. In this method, which is also known as minimally invasive pectus carinatum correction surgery (MIRPC), the chest cavity is not entered. Under general anesthesia, it is aimed to place a nickel-steel alloy metal bar presternally through a tunnel created under the skin and to correct the deformity by pressing on the sternum. At the end of 2 years, the bar is removed again under general anesthesia.

Non-Surgical Treatment

Non-surgical treatment of pectus excavatum is vacuum bell application. It consists of a silicone suction cup and a hand pump, which are produced to provide negative pressure that will create a vacuum effect on the chest wall. Successful results are obtained with its regular application in the prepubertal period, when bone development is not yet completed. (Haecker, 2011)

In the non-surgical treatment of pectus carinatum, the dynamic external compression method (orthosis/corset treatment) is frequently used in suitable patients as an alternative to surgical treatments. With a specially developed device, the anterior chest wall is kept under pressure for certain periods. The treatment is applied for 1-1.5 years and the patient is followed up at regular

intervals throughout the treatment. Determining the appropriate pressure level and the patient's compliance with the treatment are important in the success of the treatment. Compression-related pain and vasovagal symptoms, local subcutaneous hematomas and skin ulcerations are rare complications of this treatment.

Poland Syndrome And Treatment

Poland syndrome is characterized by hypoplasia or aplasia of the pectoralis muscles; It may be accompanied by anomalies such as hypoplasia (brachydactyly), finger fusion (syndactyly), breast anomalies, rib defects, absence of pectoral and axillary hair, and weakness of subcutaneous adipose tissue. It occurs in approximately one in 30 000 - 32 000 live births. (Shamberger, Welch & Upton) Generally, familial predisposition is not observed. Its etiology is not clearly known. The syndrome affects only one side of the body, 60-75% is seen on the right side.

Functional complaints are rare in patients, mostly aesthetic concerns. In addition, respiratory complaints can be seen in the presence of costal agenesis and severe chest wall defect. In patients with chest wall defects, surgical treatments should not be delayed as much as possible in order to protect vital organs such as the heart and lungs. Surgical procedures for the hand, such as syndactyly correction surgery, should also be performed as early as possible.

The treatment to be applied is determined according to the components accompanying the syndrome. The main purpose of treatment is to protect vital organs and functions. In addition, it is aimed to ensure the stability of the rib cage and to give the maximum symmetrical appearance.

Sternal Cleft And Ectopia Cordis

It is a rare congenital chest wall deformity. The sternum is formed between the 7th and 10th weeks of pregnancy as a result of the fusion of two mesenchymal leaves, starting from the manubrium and down to the xiphoid. Sternal defects occur due to complete or partial deficiency of this fusion.

In the sternal cleft, the heart, pericardium and diaphragm are in their normal localization, with normal skin cover. Cardiac defects in this group are rare and usually asymptomatic. It occurs in 1 in 50,000 to 100,000 live births. Diagnosis can be made by inspection and palpation in the neonatal period. It is very important to see paradoxical chest movements and heart movements under the skin in the diagnosis. Imaging methods are also important to detect additional malformations that may accompany.

In case of defect in the lower part of the sternum, ectopiacordis can be seen. In cases with thoracic ectopia cordis, the heart is outside the thorax without a pericardial sheath. Thoracoabdominal ectopia cordis is associated with sternal cleft, diaphragmatic defects, omphalocele, pericardial defect, and intrinsic cardiac anomalies (Cantrell pentalogy). (Gorlin & et al, 1994)

Surgical applications include different techniques from primary closure of the defect to closure with various prostheses and grafts. Surgical treatment should be performed as soon as possible after birth to prevent intrathoracic organ damage, protect it from external factors, and restore normal cardiac and pulmonary functions.

Jeune Syndrome

It was first described in 1955 by Jeune et al. It can be congenital or acquired. Acquired Jeune syndrome is a severe complication that can be seen after open pectus excavatum surgery. Congenital Jeune syndrome, asphyxiating thoracic dystrophy (ATD), is an autosomal recessive polychondrodysplasia disease. It occurs with a frequency of 1 in every 100,000 to 130,000 live births. (Kozlowski & Masel, 1976)

The thorax is narrow in the transverse and sagittal axis; It also lost its flexibility. The lungs are hypoplastic because of the narrow rib cage. Expansion of the thorax is impaired due to short horizontal ribs. As a result, alveolar hypoventilation develops due to insufficient expansion of the lungs. The morbidity and mortality of the disease vary according to the degree of lung involvement. Pneumonia and respiratory failure may occur after birth due to lung hypoplasia. Untreated babies with Jeune syndrome may die early. (Kozłowski & Masel, 1976) In moderately severe cases, recurrent pneumonia, respiratory failure and eventually ventilator dependence may occur. In mild cases, respiratory problems may decrease with advancing age. In some patients, lung functions may be found to be completely normal.

Jeune syndrome is a multisystemic disease and other accompanying skeletal anomalies, kidney failure, hepatic and biliary fibrosis, pelvic anomalies, retinal anomalies can be seen.

The main goal in the treatment of Jeune syndrome is to widen the rib cage and allow adequate lung expansion. Surgical treatment should be performed as soon as possible after delivery.

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Preventing Liver Toxicity Induced By Methotrexate

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Introduction

Methotrexate (MTX), formerly known as amethopterin, is one of several folic acid antagonists originally utilized in children with acute leukaemia (Conway and Carey, 2017; Fiehn, 2008; Martin et al., 2009; Salim et al., 2008). The data on the effectiveness of MTX derive from studies conducted in the 1960-1970s when there are no current methodological standards, but clinical experience with MTX is more than studies (Chen et al., 2009; Ginnani et al., 1992; Jolivet et al., 1983; Pınarbaşı et al., 2008). Methotrexate is also an effective drug for the treatment of adult malignancies. Toxic reactions can occur when patients use high doses of MTX, but the relationship between toxicity and concentration in adults is controversial (Yang et al., 2018). Wippel et al., in their study conducted in 2018, they investigated the safety and efficacy of high-dose MTX for osteosarcoma in young adults aged 7-38 years and difference in clearance time between the two study groups (≤ 18 years vs > 18 years) (Wippel et al., 2019). When MTX treatment is planned, pre-treatment complete blood count, erythrocyte sedimentation rate, kidney function tests (urea, creatinine), urine analysis, liver function tests (AST, ALT, GGT, ALP), hepatitis serology (HBV and HCV), HIV and tuberculosis tests must be done. Blood count, liver and kidney function tests should be followed during the treatment. Biochemical tests should be repeated once a month for the first 3 months of treatment, then every 3 months thereafter. Blood count should be applied once a week in the first 3 months of treatment and if there are no side effects, the test interval should be extended (Lindsay et al., 2009). MTX is used orally or intravenously. While it is almost completely absorbed from the gastrointestinal system at low doses, the intravenous route is preferred since the absorption of the gastrointestinal system will decrease in high doses. Oral and intravenous administration of MTX provides good tissue distribution except the central nervous system (Armağan, 2015). It reaches the maximum blood level in 45-120 minutes after oral intake. Serum half-life period is 6-8 hours. MTX is excreted in the urine rate of 65-80% within the first 12 hours by binding to albumin in the circulation. MTX accumulates in erythrocytes and hepatocytes. The clinical effect begins in 2-3 weeks, reaching a maximum in the 8th week (Güven et al., 2017; Kavala et al., 2014; Roenigk et al., 1998). Anemia, thrombocytopenia, leukopenia, tuberculosis, infections, peptic ulcer, ulcerative colitis, alcoholism, immunodeficiency, pregnancy and lactation are contraindications for MTX treatment. Excessive alcohol consumption, liver enzyme elevation, liver disease, obesity, diabetes mellitus, presence of hepatitis B and C, exposure to hepatotoxic drugs are also important risk factors for hepatotoxicity (Nast et al., 2012). The severity of the side effects of MTX is variable. The most common side effects are mild and reversible. Side effects such as nausea, vomiting, increased transaminases and stomatitis are often dose related. Generally,

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these side effects resolve after the end of treatment or dose reduction (Van et al., 2001). MTX causes side effects such as nephrotoxicity and hepatotoxicity. These side effects may occur as a result of oxidative damage caused by reactive oxygen species. There are studies in the literature stating that it should be used together with antioxidants to avoid MTX toxicity. In these studies, the occurrence of hepatotoxicity due to the use of MTX was explained as follows; the use of MTX, which is kept in the form of polyglutamate inside the cell, increases the amount of polyglutamate form in the cell and decreases folic acid levels. This causes hepatocyte necrosis (Akbulut et al., 2014; Cetin et al., 2011; Jahovic et al., 2003; Kahraman et al., 2013; Kolli et al., 2009; Kose et al., 2012; Soliman, 2009; Şentürk, 2016; Vardı et al., 2010. Approximately 30% of the patients receiving MTX treatment are discontinued due to drug toxicity (Van et al., 2001). Long-term and high-dose use of MTX, especially due to hepatotoxicity, may be interrupting the treatment or prolonging the treatment process by reducing the dose to be used. Therefore, with this study, we aimed to collect studies in the literature to prevent hepatotoxicity caused by MTX. This study was produced from my doctoral thesis, which was supported by Cukurova University Research Fund as project number TDK-2018-10773 and was finished on September 1, 2022 (decision no: 24/68-65).

Methotrexate Clinical Applications

MTX is one of the first drugs used for the treatment of rheumatoid arthritis since the 1980s. Friedman and Cronstein examined the mechanism of MTX activity in rheumatoid arthritis in their study in 2019. The reason why MTX is preferred for the treatment of rheumatoid arthritis is the explanation considered as Adenosine signal because MTX increases adenosine levels and causes adenosine to associate with its extracellular receptors, interacting intracellularly with a general anti-inflammatory effect. Studies have shown that MTX is used effective as an antagonist folate pathway by inhibiting dihydrofolate reductase when administered in very high doses (as high as 1 gram in a single dose) for leukemia, and in much lower doses (15-25 mg / week) for rheumatoid arthritis patients (Visentin et al., 2012). Rheumatoid arthritis is a common autoimmune disease that seriously endangers human health (Boone et al., 2019; Burmester et al., 2017; Pang et al., 2018). Another department where MTX treatment is dermatology. MTX is most commonly used in psoriasis in dermatology (Milojevic et al., 2001; Van et al., 2019; Thi et al., 2019). When studies investigating MTX treatment for other dermatological disorders are examined; Systemic retinoids take the first place in the treatment of Pityriasis Rubra Pilaris (PRP), which is a papulosquamous dermatosis. Dicken et al. they reported that when patients with PRP used 10-25 mg MTX a week for an average of 6 months, they had very positive results in all patients (Dicken, 1987; Lower and Baughmann, 1995; Paul et al., 1994). In addition, it has been reported that various immunosuppressives, especially systemic steroids, are used in the treatment of dermatomyositis (DM) with inflammatory myopathy and characteristic skin findings (Kasteler and Callen, 1997). In cutaneous lupus erythematosus, which is one of the cutaneous manifestations of systemic lupus erythematosus, the first choice is hydroxychloroquine, but it has been shown that it can be used in systemic steroids and other immunosuppressives in recalcitrant cases (Wenzel et al., 2005). In recent studies, it has been reported that low-dose MTX has beneficial effects on atopic eczema, which is a chronic and recurrent disease, with dry skin, itching and dyshydrotic eczema with itchy vesicles in hairless areas (Egan et al., 1994; Seyger et al., 1998). In the literature, there is only one study with MTX in Behçet's disease, which is characterized by vascular involvement such as oral and genital aphthae, ocular involvement, erythema nodosum, arthritis and thrombophlebitis

(Jorizzo et al., 1991; Lally et al., 1985; Melton et al., 1991; Torti et al., 2007). In another study by Van et al. in 2019, they evaluated the effectiveness and safety of MTX together methylprednisolone dose in the treatment of severe alopecia areata. As a result, they found that the combination of MTX and methylprednisolone was effective and safe in the treatment of alopecia areata (Van et al., 2019).

Studies to Prevent Toxicity Caused By Methotrexate

MTX inhibits mitosis of the cells by antagonizing folic acid required for deoxyribonucleic acid (DNA) synthesis of cells. MTX toxicity targets vital organs and structures of the body namely skin, GI mucosa, kidney, liver, and bone marrow. Major toxic effects of MTX, such as hepatic, renal, pulmonary, and bone marrow disorders, occur less frequently than the minor effects but may be lifethreatening. Signs and symptoms of acute MTX toxicity are based on extent and severity of organ involvement (Madke et al., 2015).

Table 1 - Studies to prevent liver toxicity caused by MTX

MTX Toxicity	Experimental animals	Human
Hepatotoxicity	Molsidomine Metformin Alpha lipoic acid Rutin trihydrate (Santa Cruz, USA) Turmeric Timokino Balanites aegyptiaca extract, Melatonin Ursodeoxycholic acid Sitagliptin Metformin Vitamin E Metallothionein Indole-3-Carbinol Inulin Tempol Glycyrrhizin acid Moringa oleifera leaf extract Coenzyme Q10 Ellagic acid Ginkgo biloba extract Acori Graminei Rhizoma	Silymarin Laurencia undulate Folic acid Leflunomide
Testicular toxicity	Carvacrol	
Nephrotoxic effect	Testicular toxicity Infliximab Lycopene Melatonin Corn oil Alpha lipoic acid Nigella sativa (black seeds) Berberine Caffeic acid phenethyl ester	Silymarin
Intestinal mucositis	Arginine, Omega 3 fatty acids nucleotides (oral formula enriched)	
Pulmonary fibrosis	Clarithromycin	
Pancreatic toxicity	Tumor necrosis factor alpha inhibitors	
Inflammation	Coconut oil	
Splen toxicity	Moringa oleifera leaf extract	
Oral mucositis	Alpha lipoic acid	
Fibrosarcoma-bearing	Vitamin c	
Over toxicity	Alpha lipoic acid	

When the literature studies on MTX toxicity in experimental animals are examined (Table 1); Daggulli et al. examined the protective effects of Carvacrol against testicular toxicity caused by MTX in rats. Twenty-four rats equally divided into three groups: group I; control treatment, group II; MTX treatment, group III; received MTX + Carvacrol treatment. On the first day of the experiment, group III received a single dose of Carvacrol, and on the second day of the experiment, groups II and III received a single dose of MTX. They found that the levels of MDA, TOS and OSI in serum and testis sampling were significantly higher in the MTX group only, compared to the control group, and significantly decreased in the MTX group compared to the control group in the TAS group. They noted that with the administration of Carvacrol, levels of MDA, TOS, OSI, and TAS were reduced relative to rats treated with MTX alone (Daggulli et al., 2014). In another study made by Aslaner et al. In 2015, they investigated the effectiveness of ozone therapy on the nephrotoxic effect of MTX. For this purpose, 18 rats is divided into three equal groups as Group 1; control, Group 2; MTX given, Group 3; MTX and ozone therapy given. Ozone therapy (25 mcg / ml) was applied to all three groups every day, and a single dose of MTX (20 mg / kg) was given to Groups 2 and 3 on the fifteenth day, and tissue and blood samples were taken at the end of the third week. Ozone therapy has been found to heal antioxidant enzymes-activated kidney tissue in rats and prevent nephrotoxicity caused by MTX (Aslaner et al., 2015). In a study conducted by Kirbas et al. in 2015, they examined the effect of anti-inflammatory infliximab active ingredient on preventing renal damage caused by MTX on rats. It was studied in three groups with 10 rats in each group. Group 1; receiving a single dose of MTX (20mg/kg), Group 2; receiving a single dose of infliximab (7mg/kg) and Group 3; receiving MTX (20mg/kg) and infliximab (7mg/kg). After five days, blood and tissue samples were taken from the rats. Consequently, at the end of five days, levels of tumor necrosis factor- α ($P=0.008$), interleukin-1 β ($P=0.036$), nitric oxide ($P<0.001$) and adenosine deaminase ($P<0.001$) were compared to Group 2 and 3. They found that it was higher, and infliximab was shown to have a strong protective effect against MTX-induced nephrotoxicity by suppressing cytokine release (Kirbas et al., 2015). In addition, in the study by Oğuz et al. examined the effects of lycopene alone or combination with melatonin on nephrotoxicity caused by MTX in rats. They divided 26 rats into four groups; Group 1; control, Group 2; given MTX (20 mg / kg), Group 3; given MTX and corn oil, Group 3 given MTX and lycopene, Group 4; MTX, lycopene and melatonin. In Group 3 and Group 4, a significant decrease was achieved in tumor necrosis factor-alpha, interleukin 1-beta and ceruloplasmin levels compared to Group 2. In addition, a significant decrease was observed in the nitric oxide level in Group 4. In histological examination, while there was damage in Group 2, it was found that there was an obvious improvement in Group 4 (Oguz et al., 2015). Çakır et al. investigated the antioxidant and anti-inflammatory effects of alpha lipoic acid (ALA) on MTX-induced kidney damage on rats. They divided thirty-two rats equally into four groups; Group 1; Control, Group 2; ALA, Group 3; MTX and Group 4; ALA+MTX. They gave a single dose of MTX (20 mg / kg) intraperitoneally to Groups 3 and 4 for kidney injury. On the sixth day, blood samples and kidney tissues received measurement of TNF- α , IL-1 β , malondialdehyde, glutathione, myeloperoxidase and sodium potassium-adenosine triphosphatase levels and histological examination results. They stated that the application of MTX caused a significant decrease in GSH and Na⁺, K⁺-ATPase activity in the tissue, and a significant increase in MDA and MPO activities. They also found that proinflammatory cytokines (TNF- α , IL- β) were significantly increased in the MTX group. They found that ALA treatment reversed all biochemical indices as well as histopathological changes induced by MTX++++ (Çakır et al., 2015). In the Medical Speciality Thesis made by Akyüz in 2015, investigated the effects of arginine, omega 3 fatty acids and an oral formula enriched with nucleotides in the MTX-induced experimental intestinal mucositis model. The study was conducted

on 32 rats by forming four groups. Group 1; Saline only by gavage was given for 5 days, Group 2; three times a day immunonutritional oral formula (5 mg/kg) was given, Group 3; After intraperitoneal single dose MTX (20mg/kg), saline by gavage was given for 5 days and Group 4; immunonutritional oral administration was given three times a day for 5 days following a single dose of MTX. Tumor necrosis factor alpha (TNF- α) and interleukin 1 beta (IL-1 β) levels were studied in serum on the 6th day, and luminol, lucigenin, glutathione, myeloperoxidase, malondialdehyde and Na⁺-K⁺ ATPase levels were studied in jejunal tissue samples. As a result, serum TNF- α and IL-1 β levels were significantly higher in the Group 3 compared to the Group 1 ($p < 0.001$). TNF- α and IL-1 β levels in the Group 4 were found to be statistically lower than the Group 3 ($p < 0.001$). In jejunal tissue, glutathione and Na⁺-K⁺ ATPase levels were lower in Group 3 than Group 1 ($p < 0.05$, $p < 0.01$, respectively), while the mean value in Group 4 was higher than in Group 3 ($p < 0.05$). In addition, tissue myeloperoxidase and malondialdehyde levels were statistically higher in Group 3 than Group 1 ($p < 0.001$, $p < 0.01$, respectively). In Group 4, myeloperoxidase and malondialdehyde levels were found to be statistically lower than Group 3 ($p < 0.001$, $p < 0.05$, respectively). Luminol and lucigenin levels in Group 3 were significantly higher than Group 1 ($p < 0.01$, $p < 0.001$, respectively); Luminol and lucigenin levels in Group 4 were significantly lower than Group 3 ($p < 0.01$). This study has shown that chemotherapy has negative effects on the small intestine mucosa, and the immunonutritional oral formula has demonstrated the protective effect of intestinal damage due to MTX (Yilmaz et al., 2018).

In a study conducted by Ahmed et al. in 2017, they examined the effect of *Nigella sativa* (black seeds) on nephrotoxicity caused by MTX in mice. There are eight mice (Swiss albino) in each group. Group 1; control, Group 2; MTX receiving (10 mg / kg), Group 3; *Nigella sativa* (black seed) receiving (0.125ml/day), Group 4; They studied with four groups receiving MTX (10 mg/kg) and *Nigella sativa* (black seed) (0.125ml/day). Malondialdehyde (MDA) and glutathione (GSH) measurements were made on the kidney homogenate on the 23rd day, and they found a increase in MDA value in Group 2 and a lower GSH level in Group 4 compared to Group 1. While in Group 2 had nephrotoxicity, no changes were found in mice in Group 4 (Ahmed et al., 2017).

In the study made by Kalemci et al. in 2018, they examined the effectiveness of clarithromycin as a protective agent in the pulmonary fibrosis model induced by MTX. They examined 30 rats as three groups; Group 1; control, Group 2; MTX (3 mg/kg) and Group 3; MTX (3 mg/kg) and clarithromycin (200 mg/kg). Histopathological changes were evaluated and normal changes were observed in the control group. In group 2, histopathological changes were found to be significantly higher in the pulmonary fibrosis MTX group compared to the other groups. As a result, they have shown that clarithromycin is effective in protecting against pulmonary fibrosis caused by MTX (Kalemci et al., 2018). In addition, Mercantepe et al. investigated the protective effects of tumor necrosis factor alpha inhibitors on pancreatic toxicity caused by MTX. They divided the rats equally and randomly into 3 groups (Group 1, healthy controls, Group 2, MTX group and Group 3, MTX+ADA group) and rats have applied intraperitoneal injection of a single dose of MTX (20 mg/44kg) in Group 2, Group 3. Group 3 received a single dose of 5 mg/kg ADA (INFLECTRA®) ip and all rats were sacrificed under anesthesia 5 days after receiving the MTX injection. Compared to those recorded in Group 1, they recorded higher edema, necrotic cell and inflammatory scores in Groups 2 and 3, edema, number of necrotic cells and inflammatory scores in Group 3, and a more pronounced decrease in Group 2. A decrease in Langerhans cell insulin and somatostatin positive interneurons islets was observed after MTX administration. They found that the islets of Langerhans and the restructuring of the pancreatic structure occurred after ADA treatment (Mercantepe et al., 2018). In the study of Rong et al., they examined that MTX

improved spinal cord injury and found that CASPASE 3, GRP 78 apoptotic factors increased in spinal cord injury, but the values decreased as MTX was administered (Rong et al., 2018). In the study conducted by Famurewa et al. in 2018, 24 rats (Group 1; control, Group 2; MTX 20 mg/kg, Group 3; MTX and 5% coconut oil and Group 4; MTX and 15% coconut oil) for the prevention of oxidative stress caused by the use of MTX with coconut oil. The parameters of SOD, CAT, GPx, GSH, MDA, Interleukin-6, C-reactive protein and nitric oxide were evaluated. They found a significant decrease in SOD, CAT, GPx and GSH values in the groups using coconut oil extract and a significant increase in Interleukin-6, C-reactive protein and nitric oxide parameters. SOD, CAT, GPx (U/mg protein), GSH (mg/g protein), MDA (nmol/mg protein) values were found respectively; in Group 1, 0.22 ± 0.01 , 9.83 ± 0.20 , 44.3 ± 1.70 , 4.50 ± 0.16 , 1.25 ± 0.30 in Group 2; 0.18 ± 0.01 , 7.46 ± 0.22 , 19.6 ± 2.00 , 3.36 ± 0.24 , 2.18 ± 0.20 in Group 3; 0.22 ± 0.01 , 8.67 ± 0.19 , 32.2 ± 0.66 , 3.91 ± 0.02 , 0.93 ± 0.23 and in Group 4; It was 0.23 ± 0.01 , 10.2 ± 0.28 , 34.5 ± 1.28 , 4.29 ± 0.10 , 1.02 ± 0.03 (Famurewaa et al., 2018).

In another study examining the protective effect of berberine against MTX-induced nephrotoxicity, the anti-apoptotic properties of berberine were demonstrated by suppressing BAX and CASPASE-3 and increasing BCL2 expression when compared to the MTX-administered group (Hassanein et al., 2019). Similarly, in another study conducted to prevent the damage of *Moringa oleifera* leaf extract to the spleen caused by MTX in mice, BAX and CASPASE 3 values were found to be higher in the MTX group compared to the *Moringa*+MTX group (Soliman, 2009).

When the literature studies with humans on MTX toxicity, Taylor et al., examined the evaluation of biomarkers of oxidative stress and apoptosis in patients with severe MTX neurotoxicity. Central nervous system therapy is an important part of ALL treatment and the most common treatment is intrathecal and high dose intravenous MTX. Treatment with MTX can cause neurotoxicity, which often leads to neurological changes, delays in treatment, and prolonged hospital stays. Oxidative stress assessment, oxidized phosphatidylcholine (PC), oxidized phosphatidylinositol (Pi) and F2 isoprostanes; have measured apoptosis with caspase 3/7 activity. They found the most consistent biomarker changes in the cases they examined, an increase in caspase 3/7 and F2 isoprostanes before acute toxicity. They found the most consistent biomarker changes in the cases they examined, an increase in caspase 3/7 and F2 isoprostanes before acute toxicity. They also found that these findings support the role of oxidative stress in MTX-induced neurotoxicity. 58 Pulmonary damage occurs as a serious adverse event in 0.3% to 8% of patients, usually during the first year of MTX treatment. Pneumonitis caused by alveolar damage, perivascular inflammation and lung infections. MTX can precipitate in renal tubules, leading to renal failure and nephrotoxic side effects due to acute tubular necrosis (Imokawa et al., 2000; Romão et al., 2014; Widemann et al., 2006). In another study investigating apoptosis in SKOV-3 ovarian cells during treatment with MTX, they noted that MTX caused an increase in BAX, BAX / BCL2 ratio and a decrease in BCL 2 ($P < 0.01$). As a result, they found that MTX caused moderate apoptosis in SKOV-3 ovarian cells (AlBasher et al., 2018).

MTX is a drug used in the treatment of various malignancies. Unfortunately, it leads to life-threatening complications, including hepatorenal toxicity. Hepatotoxicity due to drugs is one of the most common causes of liver damage (Chen et al., 2015; Lucena et al., 2008). This is because the liver is the main organ for the metabolism of many drugs or chemical agents. There are more than 1100 toxic substances that are harmful to the liver (Abboud and Kaplowitz, 2007). The increase in liver enzymes as a result of drug use is an important factor in stopping the production and distribution of the drug (Eren et al., 2004). Although severe drug-induced liver injury is an

extremely rare event, it may require orthotopic liver transplantation or result in the death of the patient (Massart et al., 2017). For example, there are studies demonstrating the protective effects on hepatorenal toxicity in other models in addition to its anticancer effects (Isoda et al., 2006; Nasri and Rafieian-Kopaei, 2014). When the literature studies with experimental animals on hepatotoxicity induced MTX, in the study made by Aşcı in 2010, examined the prophylactic effect of Misoprostol in preventing liver and kidney damage caused by MTX. In the study, 24 Wistar albino male rats were used and formed 4 groups, Group 1; Control, Group 2; MTX (20 mg/kg, ip, single dose), Group 3; MTX (20 mg/kg, ip, single dose)+Misoprostol (200µg/kg, oral, 5 days) and Group 4; Misoprostol (200µg/kg, oral, 5 days). As a result of the biochemical analysis in liver tissue, it was found a significant increase was observed in MDA levels in rats using MTX compared to the control group, while MDA decreased with the use of MP. The antioxidant enzyme CAT and SOD levels were found to be significantly decreased in Group 2, while the use of MP increased these enzyme levels. In the kidney tissue, biochemically, it was found that MTX increased the MDA level and decreased the CAT, SOD values, while MP significantly decreased the MDA level and increased the CAT and SOD values. In parallel with biochemical findings, histological examinations of liver and kidney tissues also observed findings that MTX caused oxidative damage. As a result, it was concluded that the toxic effects of MTX on rat liver and kidney tissue can be healed with Misoprostol (Aşcı and Özer, 2011). In a study by Ramachandran et al., they examined the effect of acetaminophen on mitochondrial oxidative stress, DNA and liver damage, they found an increase in AIF and BAX values (Ramachandran et al., 2011). Similarly, Çakır et al. examined the effect of alpha lipoic acid on rats in the prevention of hepatotoxicity caused by MTX. Thirty two rats is divided into four groups; Group 1; Control, Group 2; ALA, Group 3; MTX, Group 4; MTX+ALA. Groups 3 and 4 were given a single dose of MTX (20 mg / kg), while groups 2 and 4 were given intraperitoneal alpha lipoic acid for five days. On the 6th day, blood and liver tissue taken from rats were examined. TNF- α (pg / ml) and IL-1 β (pg / ml) values were found respectively in Group 1; 9.7 ± 1.3 , 10.5 ± 1.3 , in Group 3; 33.1 ± 3.5 , 31.6 ± 3.1 in Group 4; 13.9 ± 2.8 , 16.3 ± 2.7 . It was concluded that MTX caused a significant decrease in GSH and Na⁺, K⁺ ATPase activity while it caused a significant increase in MDA and MPO activity. In addition, TNF- α , IL- β values also increased significantly in the MTX group. As a result, they found that ALA treatment reversed all these biochemical indices and histopathological changes induced by MTX (Çakır et al., 2015). In another study, Erdogan et al. investigated the possible protective effect of rutin (Rutin trihydrate, Santa Cruz, USA) on MTX-induced hepatotoxicity. Twenty-two rats were divided into three experimental groups for the study; Group 1; Control-saline, Group 2; MTX, Group 3; MTX+Rutin. They took hepatic tissue for histological evaluation and biochemical analysis, analyzed the oxidative stress parameters (malondialdehyde (MDA), glutathione peroxidase (GSH-Px) and superoxide dismutase (SOD)) and liver markers in serum (aspartate aminotransferase (AST), alanine aminotransferase (ALT)). They found that there was less damage in histological examinations and increased levels of MDA, ALT, SOD and GSH-Px in the MTX+Rutin group compared to the MTX group. They found that MDA and ALT levels were higher, SOD and GSH-Px levels were lower, and AST levels were similar in the MTX group compared to the MTX + Rutin group (Erdogan et al., 2015). In another study made with experimental animals, Moghadam et al. examined the effect of turmeric against liver toxicity and oxidative stress caused by MTX. Wistar albino male rats were fed for 30 days and they were divided into 6 groups; Group 1; control, Group 2; turmeric (100mg/kg), Group 3; turmeric (200mg/kg), Group 4; MTX (20mg/kg), Group 5; turmeric (100mg/kg)+MTX (20mg/kg), Group 6; turmeric (200mg/kg)+MTX (20mg/kg). SOD, CAT and GSH-Px for cellular antioxidant defense activities and MDA for lipid peroxidation were investigated. They obtained significant signs of liver damage in the MTX group. In the

microscopic examination of liver tissue, they found that MTX treatment caused severe centrilobular degeneration, periportal degeneration, portal vein hyperemia, increased arterial inflammatory cells infiltration, necrosis, and all histopathological changes were reduced in the group given turmeric (200 mg / kg) (Moghadam et al., 2015). In addition, El-Sheikh et al. examined the effect of Timokino (TQ) (10mg / kg) on MTX-induced hepatorenal toxicity on rats orally for 10 days. For this study, four groups were designed as Group 1; control, Group 2; TQ given, Group 3; MTX given and Group 4; MTX+TQ given. They observed that caused impairment in kidney and liver functions in urea nitrogen, creatinine, alanine aminotransferase and aspartate aminotransferase in the groups given MTX (20mg/kg i.p.) at the end of the 3rd day. Moreover, they found that MTX caused a decrease in glutathione, catalase, increase in malondialdehyde levels and nitrosative stress associated with increased nitric oxide in liver and kidney. They also found that MTX caused an apoptotic effect as it lowered the caspase 3 level. When the groups given TQ at the end of the 10th day were examined, they found that MTX also reversed the inflammatory and apoptotic findings, oxidative and nitrosative stress (El-Sheikh et al., 2015). In another study by Ramachandran et al., they examined the effect of hepatitis C virus structural proteins on acetaminophen-induced liver damage in mice, and stated that there was an increase in AIF and GADD153 values compared to the control group and a decrease in GRP78 in mice with hepatitis c with liver damage (Ramachandran et al., 2015). Hamed et al. investigated the protective effect of strawberries (*fragaria ananassa*) against hepatotoxicity due to carbon tetrachloride and found that the value of BAX, CASPASE 3 decreased and BCL2 value increased in the strawberry given group (Hamed et al., 2016). Montasser et al. in 2017 examined the protective effects of balanites aegyptiaca extract, melatonin and ursodeoxycholic acid against hepatotoxicity caused by MTX in male rats. Eighty adult male rats (Sprague Dawely) weighing 190 ± 10 g were randomly divided into eight equal groups: Group 1; Control, Group 2; MTX, Group 3; melatonin, Group 4; balanites aegyptiaca extract, Group 5; ursodeoxycholic acid, Group 6; MTX + melatonin, Group 7; MTX + balanites aegyptiaca extract, Group 8; MTX + ursodeoxycholic acid. They examined liver function biomarker enzymes, liver tissue oxidative stress parameters, total antioxidant capacity, and tumor necrosis factor (TNF- α). They also performed histopathological and immunohistochemistry examinations. In the MTX group, alanine transaminase (ALT), aspartate transaminase (AST), alkaline phosphatase (ALP), gamma glutamyl transferase (GGT), total and direct bilirubin, TNF- α levels, oxidized glutathione (GSSG), malodialdehyde (MDA), nitric oxide (NO) found a significant increase. They noted that there was a significant decrease in total protein, albumin, total antioxidant capacity, glutathione (GSH), glutathione peroxidase (GPx), glutathione reductase (GR), glutathione S-transferase (GST), superoxide dismutase (SOD) and catalase (CAT) levels in the MTX-treated group. They stated that while there were positive changes in the treatment of melatonin and balanites aegyptiaca extract, no improvement was seen in ursodeoxycholic acid treatment (Montasser et al., 2017). In another experimental study, Abo-Haded et al. investigated the protective effect of Sitagliptin, a selective dipeptidyl peptidase-4 inhibitor used clinically as a novel oral anti-diabetic agent, against MTX-induced liver toxicity. They divided forty mice into four groups (10 mice each); Control, MTX, and two sitagliptin groups (pre-treated with sitagliptin 10 mg/kg/day and 20mg/kg/day) during the five days prior to MTX injection. They found that there was hydropic degeneration, apoptosis and focal necrosis in all liver regions and in biochemical analysis, there was a significant increase in serum transaminases, alkaline phosphatase and lactate dehydrogenase in the MTX group. They observed that MTX induced oxidative stress and depressed antioxidant system of hepatic tissues, activation of inducible nitric oxide synthase and increased nitrite / nitrate level in MTX group. When investigating hepatic cellular apoptosis, they found that there was an increase in Bax and caspase-3 levels and a decrease in Bcl2 levels. They

stated that all biochemical, histopathological, and immunohistochemical markers were significantly improved in the groups in sitagliptin given groups (Abo-Haded et al., 2017). In another study made in mice to prevent acetaminophen-induced hepatotoxicity, the protective effect of Folium Microcos was examined, and an increase in CASPASE3, BAX values and a decrease in BCL2 were observed in the acetaminophen given group compared to the acetaminophen+Folium Microcos group (Wu et al., 2017). In addition, Rizk et al. investigated the effect of MET (Metformin) on hepatotoxicity of MTX induced. They divided the thirty rats into 3 groups: Group 1; control, Group 2; receiving MTX treatment and Group 3; receiving MET/MTX treatment. After 7 days, they observed hepatorenal toxicity in the group receiving MTX and performed histological examinations and biochemical analysis of liver and kidney function. As a result, they observed an increase in hepatic and renal malondialdehyde levels, a significant decrease in hepatic and renal total antioxidant capacity levels and Na + / K + -ATPase in the other groups compared to the control group. They have demonstrated that MET significantly reduces hepatorenal toxicity and can prevent toxic effects on all measured parameters (Rizk et al., 2018). Amirfakhrian et al. in their study in 2018 evaluated the protective effect of vitamin E against hepatotoxicity caused by MTX using ^{99m}Tc-phytate as a radiopharmaceutical agent. They divided the rats into five groups; Group 1; control, Group 2; solvent, Group 3; Vit E (100 mg/kg), Group 4; MTX (20 mg/kg) and Group 5; Vit E+MTX. The rats continued intraperitoneal injection of Vit E for 17 days before MTX injection and for 4 days after MTX injection. ^{99m}Tc-phytate (% ID/g) in the livers, significantly increased the ID/g % level in MTX-injected rats. This study shows that ^{99m}Tc-phytate is an acceptable radiopharmaceutical agent for the evaluation of liver and spleen injuries in an animal model (Amirfakhrian et al., 2018). In another study examining the effect of vitamin E and Metallothionein in fish to prevent the toxicological effect of cadmium on the liver, an increase in CASPASE 3, AIF, BAX values and a decrease in BCL2 value were found in the group given saline compared to the group given vitamin and Metallothionein (Duan et al., 2018). In the same way, Hasan et al. in 2018 examined the therapeutic benefits of indole-3-Carbinol in adjuvant-induced arthritis and its protective effect against MTX-induced hepatic toxicity. Arthritic Sprague Dawley rats injected Complete Freund's Adjuvant (CFA) and administered Indole-3 Carbinol and/or MTX. In order to examine the anti-inflammatory and anti-arthritic effect, they examined the values of body weight, macroscopic scoring of the hind paw, level of pivotal cytokines (TNF- α , IL-6), pathogenesis, spleen index and erythrocyte sedimentation rate. They stated that both I3C and MTX by reducing inflammatory markers and mediators, alleviated inflammation and histopathological changes affecting the tibiotarsal joint. In this study, they found that I3C alleviated arthritis and liver failure induced by MTX therapy by shrinking prooxidants, regulating antioxidant defenses, and reducing pathological changes affecting the liver (Hasan et al., 2018). In another study examining the effect of coenzyme Q10 on lung and liver fibrosis in rats treated with MTX, combined therapy of CoQ10 and methotrexate was used, and blood samples for biochemical analysis, lung and liver tissue were examined for biochemical and histopathological analysis.

They found that combined treatment with CoQ10 and methotrexate resulted in improved histological picture of lung and liver tissues, modulation of liver function (elevations in serum ALT and AST and decrease in albumin), and oxidative stress biomarkers (increase in GSH level and decrease in MDA level). Autophagy genes [rapamycin mammalian target (m-TOR), Microtubule-associated proteins 1A/1B light chain 3 (MAP1LC3B) and Sequestosom 1 ubiquitin binding protein p62 (p62 / SQSTM1)] and Protein B1 (HMGB1) with simultaneous reduction in the High Mobility Group). Based on their results, they found that CoQ10 up had protective properties against methotrexate-induced lung and liver fibrosis in the current study rat model (Mohamed et al., 2018). Ebrahimi et al. examined the hepatoprotective effect of Ellagic acid (EA) against MTX-

mediated oxidative stress (OS). Thirty two Wistar rats divided into 4 groups; Group 1; control, Group 2; 10 days of oral corn oil (0.5%)+a single 20 mg/kg dose of MTX at the end of the third day (ip), Group 3; 5 mg/kg oral EA for 10 consecutive days+a single 20 mg/kg dose of MTX (ip) at the end of the third day and Group 4; 10 mg/kg oral EA for 10 consecutive days+a single 20 mg/kg dose of MTX (ip) at the end of the third day. After biochemical analysis, liver enzymes and malondialdehyde were found to be significantly higher in the MTX group. They stated that oxygen species (ROS) overproduction, decrease in mitochondrial outer membrane potential, mitochondrial swelling, cytochrome c release and increase in caspase-3/9 resulted apoptosis. Also, they noted that overexpression of proinflammatory factors, nuclear factor kappa B (NF- κ B) and interleukin 6 (IL-6) showed MTX-induced inflammation in the MTX-treated group. But they found that EA can significantly prevent OS, mitochondrial dysfunction, apoptosis, and the inflammation. Furthermore, the EA-treated rats also found that both nuclear factor erythroid 2-associated factor 2 (Nrf2) showed significant upregulation (Ebrahimi et al., 2019). Moreover, in a study of Nho et al. examined the effect of *Acori Graminei Rhizoma* on type II collagen-induced arthritis (CIA). They evaluated the water extract of *Acori Graminei Rhizoma* (WAG) in Tus, CIA mouse models. Male DBA / 1 mice were divided into five groups (NOR; n = 10, CON; n = 10, CIA + methotrexate (MTX); n = 10, CIA + 100 mg / kg WAG; n = 10, CIA + 500 mg / kg. WAG; n = 10). They analyzed hematological parameters, liver and kidney serum toxicity markers. Also, they analyzed serum levels of interleukin (IL) -6, TNF-and type II collagen IgG by enzyme-linked immunosorbent assay (ELISA). As a result, they found that WAG did not alter hematological parameters, kidney and liver toxicity markers, but could be used to treat arthritis by reducing inflammation (Nho et al., 2019). Samdanci et al. investigated the protective effect of molsidomine against hepatotoxicity caused by the use of MTX in rats. Fourty Wistar albino rats were divided into five groups as Group 1 (control), Group 2 (Molsidomine), Group 3 (MTX), Group 4 (Molsidomine-MTX) and Group 5 (MTX-Molsidomine). Average values of SOD (U/mg), CAT (k/g), GSH-Px (U/g), MDA (nmol/g), GSH (micromole/g), MPO (ng/mL) parameters was found respectively in Group 1; 0.32, 0.22, 0.83, 3.27, 2.07, 12.75 in Group 2; 0.026, 0.50, 4.69, 6.98, 2.46, 13.76 in Group 3; 0.02, 0.22, 0.96, 13.97, 1.58, 14.65 in Group 4; 0.031, 0.41, 1.52, 8.86, 2.37, 13.86 and in Group 5; 0.036, 0.32, 1.24, 11.18, 1.97, 13 (Oguz et al., 2015). Significant differences were found between groups in all parameters except MPO parameter. While GSH, GSH-Px, and GSH were significantly higher in the group receiving molsidomine and in given the first molsidomine group, the CAT level was significantly higher in the group treated with molsidomine only (Samdanci et al., 2019). In another experimental study, Ge et al. investigated the protective effect of tempol against acute hepatotoxicity caused by acetaminophen and found that it reduced pro-apoptotic protein expressions (split CASPASE 3 and BAX) and increased anti-apoptotic BCL2 (Ge et al., 2019). Kalantarive et al. examined the protective effect of inulin on MTX-induced liver toxicity in mice. In the study, 48 male mice were divided into 6 equal groups and the mice were treated orally with inulin (100, 200 and 400 mg/kg) for 9 days and given MTX (10 mg/kg, intraperitoneally) on days 7 and 9. Serum samples were taken and they evaluated the expression levels of oxidative stress biomarkers, pathological changes, apoptotic factors such as Bcl-2, caspase -3 and miR-122. The results showed that MTX induced significant liver damage and serum factors in all mice, and found a decrease in Bcl2 and an increase in caspase-3 activity and miR-122 expression compared to the control group (Kalantaria et al., 2019). Similarly, Kury et al. examined the effect of Ginkgo biloba on preventing MTX-induced liver toxicity. Rats were divided into the saline, MTX, and Gb treatment groups (Gb was administered as 60, 120, or 180 mg/kg) and the silymarin group. Overall, a seven-day protocol was adopted, in which animals received either a single daily dose of saline (with 5% DMSO) or a daily oral dose of Gb (60, 120, or 180 mg/kg) or

a daily dose of silymarin (100 mg/kg). Looking at histopathological results, they found that MTX had a significant increase in various hepatic enzymes such as alanine transaminase (ALT), aspartate transaminase (AST) and serum alkaline phosphatase (ALP). In addition, MTX contributed to the significant release of inflammatory mediators such as apoptotic (caspase-3 and c-Jun N-terminal kinases (JNK)) and tumor necrosis factor (TNF- α). As a result, they found that treatment with Gb was protective against MTX-mediated apoptosis and inflammation, as well as correcting innate antioxidative mechanisms such as glutathione (GSH) and glutathione S-transferase (GST) (Al Kury et al., 2020). In the study made by Orazizadeh et al., examining the effect of glycyrrhizin acid on BAX and BCL2 expression in hepatotoxicity caused by Titanium dioxide nanoparticles in rats, they found that there was an increase in BAX expression and a decrease in BCL2 expression (Orazizadeh et al., 2020). In the study examining the protective effects of *Moringa oleifera* leaf extract against oxidative stress and apoptosis in the liver and kidney due to MTX in mice, the BAX value was found to be higher in the MTX group compared to the group receiving *Moringa*+MTX, while the BCL2 value was found to be lower (Soliman et al., 2020). In addition, Zhang et al. investigated the protective effect of aspirin on acute liver injury due to paraquat in rats, and found that BAX, AIF and CASPASE 3 decreased, while BCL2 value increased after aspirin treatment (Zhang et al., 2020).

When the literature studies with humans on hepatotoxicity induced MTX, in meta-analysis study, over 1000 patients with osteosarcoma and acute lymphoblastic leukemia were given high doses of MTX and investigated whether the 677C> T variant was associated with the risk of liver toxicity associated with MTX therapy. All patients included in the study were evaluated for hepatotoxicity and classified according to the National Cancer Institute criteria. As a result, no relationship was found between the presence of the genetic variant in the MTHFR gene and the development of liver toxicity (Hagleitner et al., 2014). They found no relationship between the two functional MTHFR genetic variants and the occurrence of liver toxicity or other toxicities. In rheumatoid arthritis patients, MTX doses are much lower than used in cancer treatment. Generally, higher doses of MTX are thought to cause liver toxicity. Therefore, it is much more important to determine toxicity in cancer patients receiving high doses of MTX. Yang et al. found a significant correlation between the 677C> T genetic variant and the occurrence of liver toxicity in their study with adult and pediatric patients of different ethnic origins with acute lymphoblastic leukemia (Yang et al., 2012). They also classified the patients according to their ethnic origin and found that the risk of liver toxicity was high, especially in Africans and Caucasians, while Asians were not at high risk. When the age factor was evaluated, they found that the presence of liver toxicity with the 677C> T variable was more dominant in adults compared to children. However, while high doses (3.5 g / m²) of MTX are used in the treatment of childhood acute lymphoblastic leukemia, the use of lower doses (15-30 mg / m²) of MTX in adults also makes this result interesting. Certainly, the number of patients in the studies is insufficient to confirm this result. Lopez-Lopez et al. in another meta-analysis study, they investigated the relationship between the presence of 677C> T and 1298A> C genetic variants of the MTHFR gene and MTX-induced toxicity in children with acute lymphoblastic leukemia. In the pediatric population included in the study, they showed that there was no relationship between these genetic variables and MTX-induced toxicity in children, as in the study by Yang and colleagues (Lopez et al., 2013). In another study in patients with rheumatoid arthritis, Owen et al. examined 309 patients with toxicities caused by MTX treatment (Owen et al., 2013). Feinsilber et al. emphasized the importance of defining and managing toxicity in high-dose MTX administration in their study in 2018. In the literature, Lin et al., selected 1130 records with liver toxicity risk from 5903 records in 2015. All features were included in a 3-month section for classification and analyzed the records with Apache clinical Text Analysis and Knowledge

Extraction System. They emphasized the importance of classifying RA patients with MTX-induced liver damage (Lin et al., 2015). Hagag et al. investigated the role of protective silymarin on hepatic and renal toxicity caused by MTX-based chemotherapy in children with acute lymphoblastic leukemia (Hagag et al., 2016).

Conclusion

These recent studies have shown that the development of liver toxicity of MTX is associated with free oxygen radicals and hydrogen peroxide. For this purpose, melatonin, nicotinamide, methionine, vitamin E, vitamin C, n-acetylcysteine, ALA, coconut, folic acid, antioxidant agents, anti-inflammatories and vasodilator agents have been used to protect tissues from liver toxicity of MTX (Table 1). Thus, discontinuation of MTX treatments due to side effects will be prevented.

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Nutrition in the Climacteric Period

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Introduction

A woman's life consists of five stages: childhood, adolescence, sexual maturity, menopause and old age. Each of these periods shows unique physical, psychic and hormonal differences. Although each period has its own characteristics, puberty and menopause are the most important periods because of their effects on women's life (Çelik Sis & Pasinlioglu, 2013). The climacteric period, on the other hand, is the transition period between the productive and old age period of women's life, characterized by symptoms that occur as a result of changes in hormonal balance due to morphological and functional changes in the ovaries. Climacterium and menopause are often confused with each other. Menopause is a stage seen in the climacteric period. The climacteric period begins around the age of 45, which is considered the premenopausal stage, and continues until the age of 65, which is considered the old age, including the postmenopausal period. In this respect, it can be defined as the transition from the sexual maturity period to the old age period. The climacteric period; includes premenopausal, menopausal and postmenopausal periods (Görgel & Çakıroğlu, 2007).

It is stated that the average age of menopause worldwide is 51 and varies between 45-55 years. In Turkey, the median age of menopause was found to be 47 years (Özdemir & Çöl, 2004). In the 2019 data of the Turkish Statistical Institute (TUIK), it is stated that 1/7 of the female population in our country is in the menopause period (TUIK, 2019), while the life expectancy in women is 81.3 years in the TUIK 2020 data (TUIK, 2020). Based on these data, it can be said that women spend most of their lives in the menopause and postmenopausal period. The problems to be encountered for the menopausal and post-menopausal female population, which constitute a large part of the society, should be emphasized (Akdağ, Kaner, & Ayer, 2022). Age at menopause is affected by many factors such as genetic, environmental, socioeconomic and nutrition. It is emphasized that many factors such as not giving birth, smoking, high dietary fat intake and caffeine consumption accelerate the onset of menopause (Sapre & Thakur, 2014).

Menopause is not a disease but a natural and inevitable process in the life of a healthy woman (Kurt & Arslan, 2020). A healthy diet, protection and maintenance of health during and after menopause are of great importance in reducing the risks of diseases that may be caused by menopause. Chronic diseases such as obesity, cardiovascular diseases, diabetes and osteoporosis, which increase during and after menopause, can be prevented to a large extent by the combination of medical nutrition therapy and physical activity. The nutritional needs of women in the menopausal and postmenopausal period are quite different from women of childbearing age. For this reason, the individual nutrition programs of women in this period should be arranged by a dietitian, taking into account their chronic diseases and their needs specific to the menopause period (Pekcan, Şanlıer, Baş, Başoğlu, & Acar Tek, 2016).

Nutrition During Menopause

Nutrition; It is defined as the consumption of nutrients that provide the nutrients and bioactive components necessary for the continuation of life, growth and development, protection, improvement and development of health, increasing the quality of life, ensuring productivity.

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Nutrition in menopausal and postmenopausal periods is important, energy, carbohydrate, protein and fat intake, fluid consumption, vitamin and mineral intake, especially calcium and vitamin D, salt and caffeine intake are important in this period (Akdağ et al., 2022).

Energy Intake

With the menopausal period, even if the body weight remains the same, a decrease in muscle mass, an increase in fat mass and a decrease in total body water level occur. Both the absence of the menstrual cycle and the decrease in muscle mass cause the basal metabolic rate to slow down, resulting in a decrease in energy expenditure (Directorate, 2017). In addition, with the advancing age, there is a decrease in metabolic activities and physical activity levels in the body. With the decrease in physical activity levels, total energy expenditure decreases, so the prevalence of obesity in menopausal women increases (Pekcan et al., 2016). It was determined that there was an average increase of 2.1 ± 5.1 kg in body weight in the menopausal period (Guthrie, Dennerstein, & Dudley, 1999). In the American Heart Association 2013 / American College of Cardiology Obesity Guidelines, it is recommended for women to consume 1200-1500 kcal/day energy intake or 500-750 kcal/day energy deficit for body weight loss and to reduce the consumption of foods with high carbohydrate and fat content (Members et al. al., 2014). However, the recommended energy intake level according to Turkey Dietary Guidelines 2015 is 1934 kcal for a moderately active woman aged 40-49, and 1917 kcal for a moderately active woman aged 50-59 (Toplar, Kaner, & Ayer, 2022).

Carbohydrate Intake

Carbohydrates, which are the main energy source of the body, should constitute 45-60% of the total energy taken daily and at least 130 g/day of carbohydrates should be consumed. Bread, which is the most consumed grain product, should be prepared from whole grain flours or mixed whole grain flours. A diet rich in whole grains provides more dietary fiber, vitamins and minerals to the human body than refined grains. In this period, it is recommended that the bread and cereal group consume 3-3.5 portions and the pulp consumption should be 25 g/day. Therefore, at least half of the total daily grain consumption should be whole grain (Akdağ et al., 2022; TÜBER, 2015). It is emphasized that the consumption of vegetables and fruits, which are a source of complex carbohydrates, is rich in vitamins, minerals and phytochemicals, and is effective in preventing cardiovascular diseases, some types of cancer, type 2 diabetes and some eye diseases. Consuming fresh vegetables mostly raw or cooked in their own juices provides more bioavailability. In the Turkish Dietary Guidelines, it is recommended to consume 2.5 portions of vegetables and 2 portions of fruit for this period (Yabancı, Akdevelioğlu, & Rakıcıoğlu, 2012). In a study conducted in England, it was determined that women with a high habit of consuming refined pasta and rice entered menopause on average one and a half years earlier, and women with high consumption of legumes about one year later (Dunneram, Greenwood, Burley, & Cade, 2018). In another study conducted with menopausal women, it was determined that dietary lactose and galactose were not associated with the age of menopause (Rostami Dovom, Moslehi, Mirmiran, Azizi, & Ramezani Tehrani, 2019). In addition, it has been determined that simple carbohydrates increase insulin resistance, impair sex hormone binding globulin production and trigger early menopause (Dunneram et al., 2018).

Protein Intake

Consumption of quality protein in recommended amounts for age and gender is important for tissue formation, repair, and musculoskeletal health in the menopausal period, as it is in every period of life. According to Turkey Dietary Guidelines 2015, adult individuals are recommended to consume 0.8-1.0 grams of protein per body weight per day (TÜBER, 2015). It is stated that the minimum protein consumption level that should be taken to prevent lean mass loss in postmenopausal women is 0.8 g/kg per day. It is recommended that 10-20% of the daily needed energy should be met from protein sources and this ratio should be regulated according to the quality of the consumed protein source (Akdağ et al., 2022). In addition, it is recommended to consume 1.5 servings/day (120 g) of meat, chicken, fish, eggs, legumes and oilseeds group (Akdağ et al., 2022).

Osteoporosis is one of the biggest health problems that occur with the menopause period. In the Turkish Dietary Guidelines 2015, it is emphasized that high protein consumption should be avoided. In a randomized controlled study with obese women in the postmenopausal period, it was determined that medical nutrition therapy containing 1.2 g/kg/day protein was effective in reducing the loss of lean body mass. However, it has been shown that 1.2 g/kg/day protein intake causes changes in muscle cell structure and organization, and body weight loss prevents oxidative stress caused by low protein intake by regulating muscle-insulin signaling (Smith et al., 2016). Unlike this study, in a randomized controlled study with postmenopausal women, it was concluded that 1.2 g/kg/day protein intake did not support the increase in lean tissue. It is stated that more studies with a longer duration and involving more participants are needed to evaluate the effect of low or high protein intake recommendation on lean tissue gain in postmenopausal women (Rossato et al., 2017).

Oil Intake

The menopausal period is a process in which many chronic diseases occur (32). In the Turkish Dietary Guidelines, it is stated that the energy from fat should be 20-35% of the total energy need for healthy individuals. However, it is recommended that cholesterol intake be below 300 mg/day and omega-3 intake should be 250 g/day (TÜBER, 2015).

In the literature, it is thought that the type and amount of dietary fat may have effects on bone health. It has been reported that excessive dietary intake of saturated fatty acids adversely affects bone density (Wang et al., 2016). In a study on the subject, it was stated that omega-3 fatty acids affect the age of menopause. It has been determined that as oily fish consumption increases, women enter menopause later (3.3 years) (Dunneram et al., 2018). In a randomized controlled study with postmenopausal women, it was reported that 900 mg of omega-3 consumption per day significantly reduced body mass index, waist circumference, blood pressure, serum triglyceride, interleukin-6 and insulin resistance (Tardivo et al., 2015).

It is emphasized that adequate adherence to the energy-restricted Mediterranean diet is important in achieving improvements in body weight loss and metabolic health in postmenopausal women (Bajerska et al., 2018).

Vitamins and Minerals

It is emphasized that the consumption of vegetables and fruits, which are good sources of potassium and magnesium, should be increased due to the increase in the risk of chronic diseases during the menopause period. Green leafy vegetables, especially rich in folic acid, reduce the risk of coronary heart disease by preventing the increase in homocysteine levels in the blood (TÜBER, 2015).

A diet rich in antioxidant vitamins A, C and E significantly reduces serum lipid peroxide levels in postmenopausal women. For this reason, it is recommended that women in the postmenopausal period increase their intake of antioxidants such as vitamin A, beta carotene, vitamin C, vitamin E, plant flavonoids and soy isoflavones. These nutrients are generally found in fruits, vegetables, soybean, cocoa and tea leaf extracts (Ko & Kim, 2020).

Adequate intake of calcium in all age groups reduces bone mineral loss and maintains bone health. For this reason, with increasing age in women, consumption of foods with high calcium content such as milk, yogurt, cheese, green leafy vegetables should be increased for bone health. The daily calcium requirement in women is determined as 950-1000 mg/day for 19-50 years old and 950 mg/day for 50 years old and later according to Turkish Nutrition Guidelines. It is recommended to consume 3 portions of milk and products, which are sources of calcium, in the menopausal and postmenopausal periods (TÜBER, 2015).

According to the Nurses' Health Study evaluation, women with the highest dietary intake of vitamin D (528 IU/day) had a 17% lower rate of early menopause than those with the lowest (128 IU/day), and women with the highest dietary calcium intake (1246 mg/day) compared to those with the lowest consumption (556 mg/day), it was determined that the rate of entering early menopause was 13% lower (Özcan & Oskay, 2013). In a study conducted with postmenopausal

women in Korea, it was stated that vitamin D concentration should be measured regularly to maintain the optimum vitamin D level, and if the level is insufficient, individuals should be made aware of the issue and additional supplementation should be given (Lee et al., 2019).

Water and Salt Consumption

Adequate fluid intake is important for all age groups to maintain normal kidney functions, prevent urinary tract infections, kidney stones and constipation. It is very important to ensure adequate fluid consumption in the menopausal and postmenopausal periods, since the sense of thirst decreases with age. With menopause, there is a decrease in total body water as a result of frequent urination and less fluid retention in the body. For daily fluid requirement, at least two liters of water (8-10 glasses of water) should be consumed per day (TÜBER, 2015).

Since the risk of osteoporosis increases with menopause, hard water with high mineral content should be preferred. In terms of bone and dental health, it is recommended that the fluoride content of drinking and potable water should be at the level of 0.7-1.2 milligrams per liter (Akdağ et al., 2022).

In a randomized controlled study to investigate the effectiveness of increasing daily water consumption in premenopausal women on the frequency of recurrent cystitis, 140 participants were divided into a control group consuming a normal amount of water and an intervention group consuming an extra 1.5 liters of water per day. As a result of the research, it was stated that increasing water intake is beneficial in reducing the risk of recurrent cystitis in women who consume little water and have a history of frequently recurring cystitis (Hooton et al., 2018).

Salt is an essential mineral source that plays a role in the body's fluid-electrolyte balance with its sodium content. It is stated that excessive salt consumption increases urinary sodium and calcium excretion, decreases bone mineral density, and therefore increases the risk of osteoporosis. It is stated that adequate dietary calcium intake is effective in preventing osteoporosis, and the recommended amount of salt consumption affects this situation positively (Yılmaz Taşkın, Demirel, & Kumsar Karakoç, 2015). The World Health Organization recommends that adults consume 5 grams of salt per day (WHO, 2012). Excessive salt consumption during menopause is associated with many diseases, especially hypertension and osteoporosis (Öztürk & Garipağaoğlu, 2018). However, in the results of the Salt Consumption and Hypertension Community Surveys SALTURK 1 and SALTURK 2, it was stated that the average salt consumption in Turkey is 18 g/day and 15 g/day. With these studies, it has been determined that the Turkish population consumes much more salt than recommended, and it has been emphasized that applications to reduce salt consumption should be given priority (Erdem et al., 2017; Erdem et al., 2010).

Conclusion and Recommendations

As in every period of life, a healthy diet, protection and maintenance of health in the menopause and postmenopausal period are of great importance in reducing the risks of diseases that may be caused by menopause. During the climacteric period, the nutritional needs of women differ from previous years. In this context, the energy, macro and micro nutrient needs of the individual should be provided with individually planned nutrition programs. Individual nutritional recommendations should be made, taking into account the symptoms experienced by women in this period, hormonal changes and their needs specific to the menopause period.

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A Current Practice in Dentistry; Endocrowns

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Introduction

Post-supported basic restorations and crowns are most commonly used in the restoration of teeth with root canal treatment and excessive tissue loss (Dietschi & et al., 2008; Zarow, Devoto & Saracinelli, 2009; Ma & et al., 2009). Despite the fact that some clinical success was achieved with the use of intracanal posts, the disadvantages of this are: it causes loss of healthy tissue in the tooth, requires additional processing and time, and causes fractures in the tooth root (Lazari & et al., 2013). Pissis proposed a new technique in which a pulp chamber was used for the retention of crowns without the need for posts. Bindl and Mormann defined “endocrowns” as adhesive crowns for the purpose of restoring root canal-treated posterior teeth that have complete coronal hard tissue loss (Bindl & Mormann, 1999).

Thanks to developments in advehise dentistry, restoration of teeth that have undergone treatment (such as root canal treatment) can be done more conservatively by using endocrowns. Endocrowns combine the crown, core, and intraradicular post in a single component; therefore, they represent monoblock restorations. Endocrown restorations, unlike traditional approaches such as using intraradicular posts, are fixed to the interior of the pulp chamber and to the cavity margins. This results in a micro and macro-mechanical retention, as provided by the adhesive cementation as well as pulp walls (Cunha & et al., 2017).

Developments in computer-aided design and manufacturing (CAD/CAM) have provided more accurate, aesthetically pleasing, and less time-intensive restorative procedures for both material processing and tooth restoration. The different materials that are made available with this technology give clinicians the opportunity to choose the most suitable materials for them.6 In the literature, for glass ceramics that are reinforced with lithium disilicate or leucite, it has been found that feldspathic glass ceramics and resin composites are the best choices for endocrown production, as they show higher flexural strength and can withstand occlusal forces during chewing (Otto, 2004; Biacchi & Basting, 2012; Bindl, Richter & Mörmann, 2005; Zhu & et al., 2017). Resin-based CAD/CAM blocks developed in recent years constitute an important alternative to traditional ceramic blocks. The mechanical properties of resin-based materials and their abrasive behavior on enamel provide significant advantages over glass ceramics, which cause severe abrasion on the enamel of fragile and opposing teeth (Lin & et al., 2010). This review’s purpose is to examine the pros and cons of endocrown restorations and to evaluate current materials and studies.

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Endocrowns

Alıntılar metin içinde ve parantez kullanılarak verilmelidir. Metin içi atıflar tek yazarlı eserlerde (Anaz, 2014), iki yazarlı eserlerde (Bekir & Ulaş, 2003), üç yazarlı eserlerde (Kul, Faruk & Deniz, 2016), üçten fazla yazarlı eserlerde ise (Gerekvar & ark., 2018) şeklinde olmalıdır. Bir yazarın aynı yıl içerisinde yayınlanan eserlerine atıf yapılırken (Yılmaz, 2014a, 2014b) şeklinde verilmelidir.

Deciding on the indications for root canal-treated teeth with excessive dental tissue loss is difficult, as there are many restorative materials and restoration options. It has been stated that the main reason for the extraction of these teeth is prosthetic, periodontal, and endodontic failures. With advances in dentistry, a more conservative approach is recommended to restore root canal-treated teeth using endocrowns. (Figure 1A) (Olçay, Ataoglu & Belli, 2018).

Endocrown restorations can be applied to all teeth on the arch, and they are generally a better recommendation for molars. Premolars and anterior teeth, meanwhile, have a small pulp chamber and are subject to lateral forces of larger magnitudes (Sedrez-Porto & et al., 2016; Rayyan & et al., 2019; Al-Dabbagh, 2020). Endocrowns are also indicated in cases of excessive coronal tissue loss and limited interocclusal space when teeth have short, obliterated, dilated, or curved roots (Biacchi, Mello & Basting, 2013). The performance of endocrowns is affected by many factors, including material types, loading axis, and preparation patterns. (El Ghouli & et al., 2019; Altier & et al., 2018; Taha & et al., 2018).

Advantages of endocrowns

- Endocrowns are a conservative technique.
- They require a much shorter session time.
- The tooth/restoration interface of the restored teeth is better at evenly distributing chewing forces across the tooth structure.
- It can be applied to teeth with curved roots and calcified canals.
- This can be applied in cases where the interocclusal distance is insufficient.
- Healthy tooth tissue is preserved.

Disadvantages of endocrowns

- Endocrowns are unsuccessful in the following cases; if adhesion is not achieved or the depth of the pulp chamber is measured below 3 mm, the thickness of the peripheral walls must be thicker than 2 mm.
- Endocrowns can be contraindicated if there is improper occlusion (parafunction).
- In tooth preparation, a 1.5-2 mm ferrule should be made to prevent fractures in the tooth (Singh & et al; 2019).

Classification of endocrown restorations

The classification of endocrown restorations is based on the amount of tooth tissue that remains after preparation.

1. A tooth preparation with a minimum of two solid walls is described.
2. A tooth preparation is described in which a maximum wall has a height of $>1/2$ of its original measured height.

3. A tooth preparation is described in which all of the walls are $<1/2$ of their original height (Belleflamme & et al., 2017).

Construction Stages of Endocrown

Cavity Preparation

The purpose of this preparation is to obtain a large, stable surface that resists the compressive stresses known to occur in molars (Zogheib & et al., 2011). The surface was prepared parallel to the occlusal plane. This provides resistance to stress along the major axis of the tooth (Fages & Bennisar, 2013). As a result, lower stress levels are achieved compared to teeth with prosthetic crowns (Dejak & Miotkowski, 2013).

Thanks to developments in adhesive cementation systems, the need for macroretentive preparation has decreased for crowns (Lander & Dietschi, 2008). The pulp chamber space also provides stability and retention. The trapezoidal shape of the mandibular molars, as well as the triangular shape of the maxillary molars, help increase the stability of the restoration. In addition, additional preparation isn't required. The saddle form of the pulp base helps increase stability. Due to the adhesive properties of bonding materials, this anatomy makes the use of root canals redundant (Figure 1B).

The quality of the endodontic treatment was evaluated from radiography before starting the endocrown preparation. Before dehydration of the teeth, the color of the material to be used was selected. The existing restorations were removed. Attention is paid to the width of the coronary chamber and the amount of enamel remaining on the edges. It is used with diamond burs to provide sufficient space (about 1.5 mm) for restoration. A diamond bur was used to obtain the proper surface on the mesial and distal walls of the tooth and to remove unsupported enamel. The cavity was finished with the same diamond burs used during the low-speed rotating preparation. All interior walls are rounded to facilitate impression-taking, restoration placement, and removal (Cunha & et al., 2017).

As deep subgingival carious lesions and restorations deepen, the gingival marginal enamel keeps thinning down until it reaches the cement-enamel junction (CEJ), and at this point, bonding to the enamel is impossible. Margins that are placed apically to the cement-enamel junction on dentin are known to be more prone to microleakage caused by differences in thermal expansion between the tooth structure and restorative material and their respective thermal expansion coefficients. Incomplete hybridization between the adhesive system and collagen fibrils is also a cause of microleakage (Stockton & Tsang, 2007).

In cavities containing subgingival margin, a technique known as deep margin elevation, proximal box elevation, or relocating cervical margin, also known as opensandwich technique is used. This technique utilizes a direct restoration approach, placed solely in the deep apical portion, during preparation to raise the cavity margin to a more favorable and coronal position for the final part of restoration fabrication and cementation. The oral environment is exposed to direct restoration by deep margin elevation. This extra interface of direct restoration can potentially leak, and there are some concerns that it may be associated with the increased failure rate with this technique (Frankenberger & et al., 2013).

Vertolli et al. showed that deep margin elevation results in a reduction in the fracture of ceramic restorations when the cavity margins extend below the cement–enamel junction. Marginal integrity has been preserved in restorations, both direct and indirect, by using the deep margin elevation technique. This suggests that deep margin elevation may be a viable treatment option, as restorations are made difficult by subgingival margins. No difference was found between the materials (resin-based composite, glass ionomer, resin-modified glass ionomer, and bulk-fill composite) used for proximal box elevation (PBE) in terms of margin quality or fracture resistance.

It is stated in the literature that any of the resin-modified glass ionomer, composite resin, or glass ionomer materials may be suitable for PBE procedures (Vertolli & et al., 2020).

Material Selection

In studies on endocrowns, nanoceramic resin, reinforced composite resin, lithium disilicate glass-ceramic, zirconium reinforced lithium silicate, feldspathic ceramic, and hybrid ceramic blocks were used. For the purposes of endocrown restorations, the most widely used material is lithium disilicate glass-ceramic (LDS), as its biomechanical properties fit. (Kuijper & et al., 2020; Zhu & et al., 2020; Einhorn & et al., 2019).

CAD-CAM materials used in endocrown restorations: Vita Suprinity (VS; VITA Zahnfabrik, Germany), IPS e.max CAD (EMX; Ivoclar Vivadent AG), Vita Enamic (VE; VITA Zahnfabrik), Grandio blocks (GR; VOCO), and Lava Ultimate (LU; 3M ESPE) were investigated using Finite Element Analysis (FEA) and Weibull Analysis (Hidaka & et al., 1999; Gonzaga & et al., 2011). As a result of these analyses, the LU and GR models showed a more balanced distribution of stress. Weibull analysis has also revealed that the 5 models performed similarly while under normal occlusal forces. During clenching, the VE and LE models achieved the highest chances. The fracture strength of GR is significantly higher than that of other materials (Zheng & et al., 2021).

In recent years, continued research on biocompatible materials that have mechanical and physical properties that are similar to those of natural dental tissues has produced a new generation of resin-based ceramic restorative materials. The hybrid nature of resin-based ceramic blocks provides a measure of elastic modulus (12.8 Gpa) that is similar to dentin. The advantages of resin-based ceramic restorations are that they show better fracture resistance and less crack propagation than some CAD/CAM ceramics (El-Damanhoury, Haj-Ali & Platt, 2014; Chen & et al., 2014; Magne & et al., 2010). Today, CAD/CAM blocks that are resin-based with different contents (Vita Enamic, Lava Ultimate, Brilliant Crios, Cerasmart, Shofu Block, and Grandio Block) are available to physicians.

Cementation

Restoration prepared using different dental design programs was produced with the help of milling devices (Figure 1C). The rod area of the restoration produced in the digital system is cut with the help of a diamond bur and removed. The endocrown is then placed in the patient's mouth to assess interproximal contact and marginal integrity. Occlusion control of the restoration was performed. Corrects sections that prevent occlusion restoration are repositioned and marked. When evenly distributed occlusal contacts are observed, polishing is performed with a diamond-containing tire rotating at low pressure and low speed. Ceramic blocks reinforced with glass ceramic, lithium silicate, and zirconia are sintered according to the instructions of the manufacturer (except resin-based CAD/CAM block). Pigments and glaze materials are used to characterize and polish the restoration. Resin-based blocks are only polished with a rubber diamond polisher.

Before cementation of the endocrown, the cavity and adjacent teeth were isolated. Restoration surfaces produced from ceramic-containing blocks are roughened for 20 seconds with hydrofluoric acid and then rinsed for 20 seconds before leaving them to dry. The inner surface of the block, which is produced from resin-based blocks, is roughened by sandblasting. Adjacent teeth are protected by polyester tape; enamel is roughened with phosphoric acid to increase retention and washed.

Dual-cure resin cement was prepared according to the instructions of the manufacturer and applied to the tooth surface. The restoration was then placed on the prepared tooth surface. Excess cement on the proximal surfaces was removed with dental floss and with an explorer on lingual as well as buccal surfaces and polymerized (Cunha & et al., 2017) (Figure 1D).

In vitro studies on endocrown restorations

Many studies have compared the mechanical and physical properties of endocrowns. When Chang et al. compared the failure modes and fracture resistance of endocrowns to conventional post-core supported crowns in maxillary premolars, they found that the endocrowns showed higher fracture resistance compared to conventional crowns; however, between the two groups, there was no significant difference in terms of failure modes (Chang & et al., 2009).

El-Damanny et al., in an in vitro study in which they examined the fracture resistance and marginal leakage concerning ceramic endocrowns produced using 3 different CAD/CAM made of lithium disilicate, resin nano-ceramic, and feldspathic porcelain on maxillary molars. It has been shown that resin nanoceramic endocrowns have noticeably greater fracture resistance (El-Damanny, Haj-Ali & Platt, 2014).

An in vitro study concerning mandibular molars found that when the fracture strength with respect to ceramic endocrowns strengthened with lithium disilicate and traditional crowns supported by glass fiber posts was examined compared to conventional crowns, endocrowns showed higher fracture strength. Furthermore, the failure pattern for both groups was similar. In addition, it is characterized by tooth fracture related to restoration displacement (Biacchi & Basting, 2012).

A study conducted on adhesive restorations of endodontically treated anterior teeth found that failure modes and fracture resistance concerning endodontically treated upper incisors that had been restored with endocrowns made of ceramic or composite blocks were examined. No significant difference was observed (Ramirez-Sebastia & et al., 2014).

Taha et al. Conducted an in vitro study to evaluate the effect that occurs when changing the margin designs concerning endodontically treated teeth that have been restored using polymer-infiltrated ceramic by the method of endocrown restorations on fracture resistance; it has been observed that endocrowns that have shoulder finish and axial reduction lines have greater fracture resistance (Taha & et al., 2018).

In the literature, studies evaluating the biomechanical behavior of teeth that have been endodontically treated and restored by utilizing different endocrown extensions in the pulp chamber have concluded that greater elongation of the endocrowns results in greater mechanical performance (Dartora & et al., 2018; Silva-Sousa & et al., 2017).

Altier et al. reviewed the fracture resistance of three endocrowns to compare them. They were made of two different indirect resin composites and lithium disilicate ceramics. Finally, they determined that, compared to the indirect composite groups, lithium disilicate ceramic endocrowns showed a higher fracture strength. (Altier & et al., 2018).

A study by Darwish et al. exhibited that endodontically treated maxillary premolars that were restored using resin nanoceramic endocrowns exhibited superior internal adaptation when compared with endocrowns that were restored using lithium disilicate. Finally, the endocrown preparation with smaller axial wall divergence ("G" grade) provided a better internal fit (Darwish, Morsi & El Dimeery, 2017).

In the study by Anton et al. fifty extracted mandibular molars were used; it has been reported that the fatigue resistance concerning endodontically treated molars restored using fiber-reinforced endocrown restorations does not increase, and the potential effect of fiber supports may be reduced due to the high thickness of the restorations. On the other hand, it has been observed that fiber reinforcement positively affects the failure mode and crack propagation after fatigue failure, thus increasing the chance of reparability (Anton & et al., 2021).

In vivo studies on endocrown restorations

Sedrez-Porto et al. evaluated clinical trials. They compared endocrown restorations to conventional treatments while utilizing direct composite resin, intraradicular posts, or onlay and inlay restorations. Endocrowns have been reported to be an improvement over conventional methods, and in the worst cases performed similarly (Sedrez-Porto & et al., 2016).

Bindl and Mörmann have also reported that 19 adhesive-bonded CEREC endocrowns (15 molars and 4 premolars) operated clinically successfully for 28 months, while only one of the endocrowns failed due to recurrent caries (Bindl & Mormann, 1999).

Belleflamme et al. reported that despite the presence of occlusal risk factors, such as extensive coronal tissue loss, adverse occlusal relationships, or bruxism, endocrowns can still be reliable as an approach to restoring severe cases of damaged premolars and molars (Belleflamme & et al., 2017).

Otto et al. concluded that the success rates of feldspathic endocrowns in premolars and molars were 75 and 90.5, respectively 75%, and 90.5% over a 12-year follow-up. In another clinical study, endocrowns showed a clinical success rate ranging from 94% to 100% in 3 years, while the 10-year success rate was 98.8% (Otto & Mörmann, 2015).

Conclusion

Endocrowns have superior fracture strength compared to teeth restored with post-core retaining crowns. Therefore, it represents a promising treatment alternative for root canal-treated molars, preserves the structure of the tooth, and is suitable for minimally invasive dentistry.

Four essential factors explain the long-lasting and successful effects of endocrowns: correct selection of restorative materials as well as correct preparations of the tooth. In addition, the location of the restoration margin and the correct case selection are important factors.

Although feldspathic and lithium disilicate ceramic blocks were initially used in the treatment of endocrown restorations, resin-containing blocks have recently been used for endocrown restorations.

Despite the gains in popularity of endocrowns, it is doubtful whether this restoration method could replace conventional restorative treatments that utilize the post-core in teeth treated with the anterior root canal. To the extent of our knowledge, laboratory and clinical evidence proving the safety of endocrowns for use in anterior teeth is lacking. Therefore, more evidence is needed, supported by long-term clinical trials.

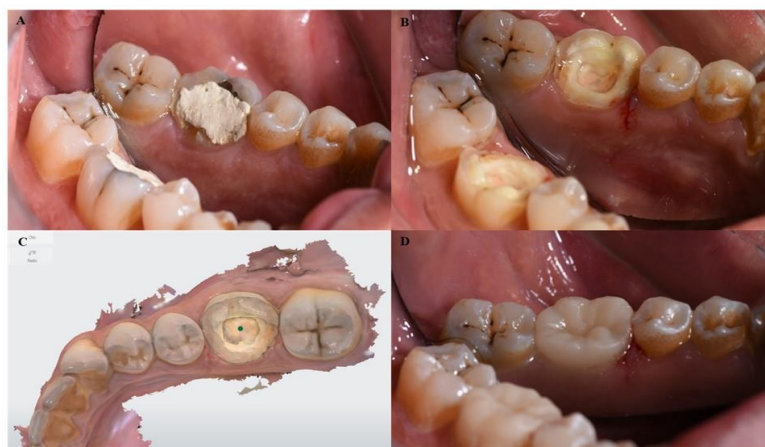


Figure 1A. Intraoral view of tooth before treatment, **B.** Cavity preparation of tooth **C.** Intraoral scanning of the cavity, **D.** Intraoral view of the completed restoration

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Ocular Delivery Systems and Applications Developed by Qbd Perspective: From the Past to the Future

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1. INTRODUCTION

Eye is one of the most complex organs that converts the light reflected from objects into electrical impulses and transmits them to the brain via the optic nerve. The eye might be investigated in two different perspectives which are anatomical and functional. In context with anatomical. The eye composes two different parts; anterior and posterior segments. While the anterior segment consists of the cornea, anterior chamber, iris, posterior chamber, ciliary body and lens, posterior segment consists of the vitreous, retina, retinal pigment epithelium, and choroid (Figure 1).

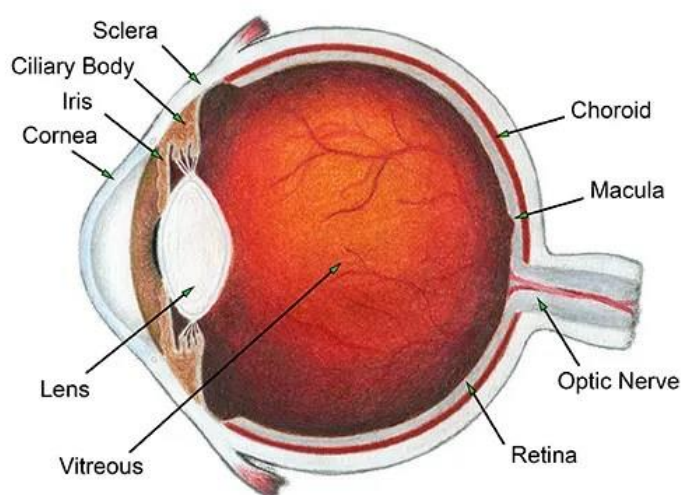


Figure 1. Anatomical investigation of eye division (Pautler, 2021)

The major duty of the components in the anterior segment is permeating the light stimuli to the posterior segment as well as that focus on the macula. The posterior segment, which forms the posterior 2/3 of the eye, includes the vitreous humor, retina, macula, and optic nerves (Gaudana, Jwala, Boddu, & Mitra, 2009; Wadhwa, Paliwal, Paliwal, & Vyas, 2009). In addition, the

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eyeball can be functionally examined in three layers. The outermost protective layer consists of the sclera and cornea. The cornea is a transparent structure with a diameter of 11.7 mm and a thickness of 0.5 - 0.7 mm, which protects the eye against infections, physical damage, and transmits light to the lens and retina by refracting it (Yasukawa et al., 2004). The sclera protects the eye from internal and external forces and ensures that the shape of the eye remains intact is the structure of the uttermost layer of the eye. The middle layer includes the ciliary body, choroid, and iris. The choroid is a highly vascularized structure located among the retina and the sclera. The iris is the structure that plays a role in regulating the amount of light reaching the retina by controlling the pupil opening (DelMonte & Kim, 2011). The retina is the layer that forms the inner layer and contains the basic nerve cells and structures for the sense of sight. And last structure is the ciliary body controls the shape of the lens (Müller, Pels, & Vrensen, 2001).

1.1 Ocular Barriers

The ocular drug delivery process is difficult due to the anatomical and physiological barriers in the eye. In the treatment of superficial diseases of the eye, systemic drug administration is not a preferable option as so the obstacle components such as the blood-aqueous barrier and the blood-retina barrier block desired level of the drug (Figure 2). Hence, for overcoming of this problem, high systemic drug dosage is being applied; unfortunately, this approach causes side effects in patients. For this reason, topical drug administration is frequently preferred in the treatment of superficial diseases of the eye, as well as eye drops constitute more than half of ophthalmic preparations (Singh, Ahmad, & Heming, 2011).

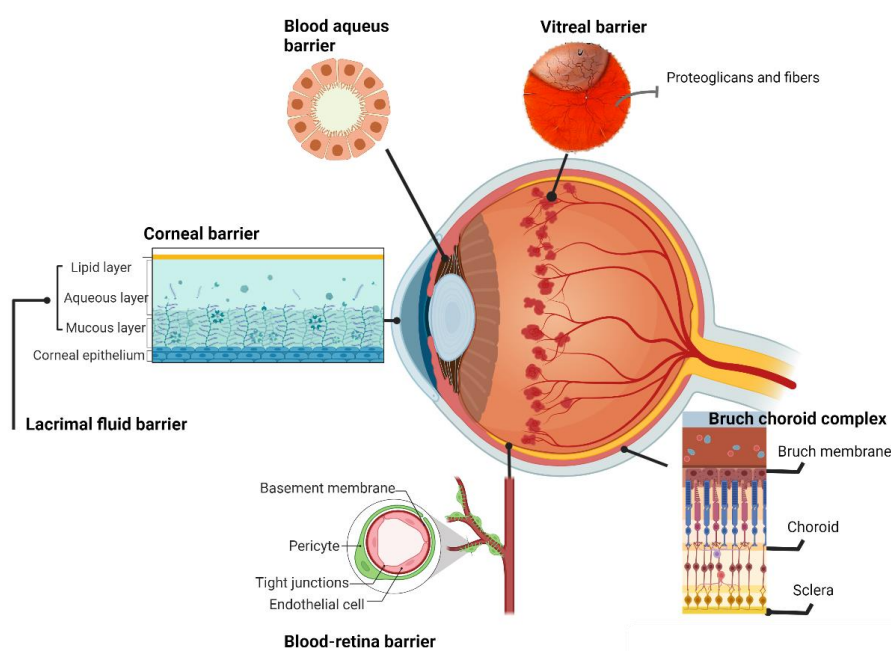


Figure 2: Encountering the barriers on ophthalmic administration

However, low ocular bioavailability (< 5%) is observed with topical administration of drugs to the eye. The reasons for this can be summarized as follows (Kompella, Kadam, & Lee, 2010):

- After the drug is applied to the eye, the drug solution is diluted with lacrimal fluid.
- The volume of lacrimal fluid in the eye is approximately 7 to 10 μL , as well as the volume of each drop applied to the eye is approximately 50 μL . Therefore, approximately 80% of the

amount of drug administered immediately protrudes from the surface of the eye or is removed from the conjunctival tract.

- Topically applied drug continues to dilute with increasing tear, normal tear cycle as well.
- Consequently, regarding other barriers in the eye, the penetration of the drug into the back of the eye is restricted.

Solutions are one of the traditional ophthalmic drug forms that are used ubiquitously forms for the topical application of drugs to the eye. The crucial points regarding the solution's formulation are solubility, ocular toxicity, pKa, the effect of pH, tonicity, buffer capacity, viscosity, compatibility with other ingredients in the formulation, preservatives to be used, comfort in eye application and can be listed as convenience (Gericke et al., 2019; Winter, Anderson, & Braun, 2010).

In order to boost the ocular bioavailability by decreasing the loss of the drug, dozens of strategies (such as increasing viscosity, developing into the ointment, adding a cationic agent, prodrug development, and penetration enhancer addition) had been utilized by the researchers (Gaudana, Ananthula, Parenky, & Mitra, 2010). Unfortunately, these strategies were not very effective at improving bioavailability.

1.2 Novel ocular drug delivery systems

New drug delivery systems are being developed since traditional strategies are not adequately efficient in terms of increasing the accommodation in the precorneal area, preventing it from moving away from this area, and increasing drug penetration through the cornea. These systems can be listed as polymeric nanoparticles, solid lipid nanoparticles, nanoemulsions, nanosuspensions, liposomes, micelles, dendrimers, hydrophilic gels, inserts, and ocular minitables (Hornof, Toropainen, & Urtti, 2005; Irimia et al., 2018; Urtti, 2006)

The pharmaceutical industry is encountered many problems for instance insufficient bioavailability and low water solubility of newly developed drug molecules. Hereby, the requirement is available to occur the action taken promptly for improving the novel drug delivery system that overcomes these disadvantages (Urtti, 2006). These delivery systems should not have acute and chronic toxicity, as well as they should have adequate drug loading capacity, drug targeting, and controlled release properties. Meanwhile, the chemical and physical stability of the drug must be ensured. Size reduction is not one of the best methods only, it is also increasing the solubility of poorly water-soluble agents of biopharmaceutical classification system (BCS) classes II and IV (Freitas & Müller, 1999; Fresta, Guccione, Beccari, Furneri, & Puglisi, 2002).

Colloidal systems are drug delivery systems in micron and nanometer sizes, in solid or semi-solid form and show all the advantages of controlled release systems (Yalpani, 1997). In addition, they increase solubility and bioavailability of inferior drugs by acting as preservatives for sensitive drug compounds (Fialho & da Silva-Cunha, 2004). Parameters that need to be controlled to select the most suitable colloidal system are indicated in the **Table 1** (Chrysanthia Freitas & Müller, 1998; Müller et al., 2001; Winter et al., 2010).

Table 1: Specifications that affecting on the colloidal systems (Chrysanthia Freitas & Müller, 1998; Müller et al., 2001; Winter et al., 2010)

Parameters of colloidal particles				
Electrostatic	Morphological	Solution	Optical	Quantitative
Surface charge	Size	Turbidity	Refractive index	Loading capacity
Zeta Potential	Molecular weight	viscosity stability	Coefficient absorption	Encapsulation efficiency

Moreover, regarding the carrier systems: *in-vivo* degradation time, toxicity, biocompatibility with ocular tissues, and excretion time from the body are also important except for the specified parameters in the Table 1.

2. QUALITY BY DESIGN (QBD) – OCULAR APPLICATIONS

Quality by Design (QbD) is a design concept that aims to cover the entire lifecycle of a product, from the design stage to the marketed stage, as well as an initiative initiated by the United States Food and Drug Administration (FDA). QbD is a new paradigm that emphasizes understanding the product and process within the product development process and throughout the product lifecycle. As can be seen from the studies in the literature, the basis of QbD studies is the scientific understanding of the product and the process, hereby statistical experimental designs (Design of Experiments, DoE).

In the literature, reducing the number of experiments and obtaining more useful data. Numerous experimental designs have been described that can provide. If the purpose of an experimental design is to classify many variables as “dependent” and “independent”, first-order designs such as Plackett-Burman or Taguchi may be preferred.

However, if the aim is to predict a response variable mathematically (estimation) or to optimize a process, quadratic models such as Box-Behnken, Central Composite, or Optimal designs should be preferred. As seen in the literature examples, the use of appropriate software in the creation is a crucial part of the process as much as selecting the suitable type of the DoE model, and the quality of the data to be obtained from the study as well. By looking at the characteristics given in **Table 1** (dependent), the roadmap used in the selection of the appropriate formulation can be determined. Determining of the levels of independent variables in two-level experimental designs is a critical point. If the gap between the low and high levels is too narrow, a significant factor may be considered meaningless, or on the contrary, if the gap between the two levels is too wide, a nonsignificant factor may be considered significant under normal conditions and may mask the effects of other factors (Ahuja, Ferreira, & Moreira, 2004; Tekade & Chougule, 2013).

Khan et al., investigated the main effects of various adjuvants and process parameters using a seven-factor, twelve-experiment Plackett-Burman design and used X-Stat® software. For this purpose, cumulative drug release was determined as the independent variable, and the lag time and the time when 50% of the total drug was released were considered by constraints. In the quantitative analysis, the most significant factors were determined, and the significance of these factors was tested by analysis of variance (Khan et al., 2000).

Kiss et al. determined the changing active substance (w/w, %), lipid (w/w, %), and surfactant (w/w, %) ratios; investigated their effects on particle size (nm), zeta potential (mV),

loading capacity (%EE) and PDI using two-factor, eight-experimental full factorial design. Consequently, it was observed that the surfactant ratio had a significant effect on both size and EE% ($p < 0.05$), it was determined that the lipid ratio did not have a significant effect ($p > 0.05$) (Kiss et al., 2020)

The effect of formulation components on the release of besifloxacin- loaded nanostructured lipid carrier was studied using Design-Expert® software (Stat-Ease Inc., USA) with the aid of a Box-Behnken Design by using amount of drug (X_1), lipid:polymer ratio (w/w, %) (X_2) and surfactant concentration (X_3) are independent variants, as well as the particle size (Y_1) and %EE (Y_2) are dependent variants. ANOVA was used to validate the optimization design and correlation of response of 3D plots was generated. As a result of the analysis of variance, it was seen that the factors had significant effects on the response variables ($p < 0.05$) (Ferreira et al., 2007).

Fangueiro et al. used Softisan® 100 (w/w, %), Lipoid S75 (soybean phosphatidylcholine) (w/w, %), and Poloxamer 188 (w/w, %) to had a lipid nanoparticle with a water/oil/water structure containing positive potential. In the Taguchi factorial design model performed with CTAB, it has been shown that the lecithin concentration in the formulation (Lipoid S75) is the key factor affecting the nanoparticle size and PDI ($p < 0.05$). The optimized formulation containing CTAB (0.5% of the lipid phase) appeared to be biocompatible in the human retinoblastoma cell line (Fangueiro et al., 2014).

Recently, formulations can be created by using 2 different design models at the same time in order to obtain precise results. Paclitaxel nanoparticles were prepared by Yerlikaya et al. using both Plackett-Burman and Box-Behnken designs. After determining paclitaxel amount (w/w,%), PLGA amount (w/w,%), PLGA molecular weight, PLGA terminal group, surfactant concentration (w/w,%), surfactant type, homogenization rate (rpm) and homogenization time parameters (min) as potential risk factors, these risk factors effects were specified on the critical quality properties of nanoparticles, which were determined using the Plackett-Burman experimental design. As a result, it was seen that the amount of PLGA (w/w,%), surfactant concentration (w/w,%), and homogenization rate (rpm) of the emulsion affected the particle size, zeta potential and encapsulation efficiency properties of paclitaxel nanoparticles, therefore these factors were critical formulation and process parameters. The formulation and process parameters were investigated in Box-Behnken experimental design after determining critical factors as a result of the Plackett-Burman experimental design.

Fifteen formulations in the Box-Behnken test matrix were prepared and analyzed for particle size, zeta potential, and %EE. When the results obtained were evaluated by regression and analysis of variance, it was seen that the mathematical models established could explain the effect of factors on particle size and encapsulation efficiency ($p < 0.05$), but the model established for zeta potential was not statistically significant ($p > 0.05$) (Yerlikaya et al., 2013).

4. FUTURE SIGHT

Process validation approaches consisting of different stages adopted in drug production for the last 10 years have brought significant advantages in the production of a targeted quality drug product. Although QbD-based product development, design, and continuous process verification phases require detailed planning and investment, it has been the most beneficial approach in terms of stakeholders (patient, industry, official authority) that concern patient health in general.

For the design and development of desired robust ocular products in terms of quality target product profile (QTTP) is mandatory given the biopharmaceutical properties of the active substance. These properties are called critical quality properties (CQAs) and include physical (particle size distribution/morphology, polymorphism, water solubility), chemical (pKa, chemical,

photolytic, and oxidative stability), biological/microbiological property (partition coefficient, membrane permeability, bioavailability, microbial limits) or the property of an output material, including the finished drug product, that must be within an appropriate limit to achieve the desired product quality.

The criticality of an attribute is decided by the degree of damage done to it. The ocular product design defines whether the product meets the patient's requirements and maintains its performance throughout the shelf life specified both by clinical and stability studies. It is obvious that it provides benefits both on the basis of the patient and company while reducing the possible errors in production to the lowest levels and ensuring the delivery of quality products to the patient. However, since the application of current approaches in process validation requires a significant culture change; regulators, industry, and academics, all together with a multidisciplinary approach needs to work.

5. CONCLUSION

DoE approach provides the most appropriate result by changing many variables at once, instead of changing one variable at a time, which is the traditional method. In this way, many factors are evaluated both simultaneously and independently of each other. Thus, the interactions between the variables are more detailed, interpretable, and clear in a short time. Based on these studies, it is clear that the designs of nanoparticles have been prepared successfully with the QbD perspective. In this way, in the future, it will be possible to use drugs that are clinically critical and costly more effectively and to improve new drug carrier systems in a less complicated and time-consuming manner.

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Vaginal Infections

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Vaginitis term has a wide definition that is including infectious, inflammatory disorders and substitutions in the normal vaginal flora. Vaginal infections topic will be discussed in this chapter particularly.

General Epidemiology

Epidemiologic studies usually fail to define the real prevalence of vaginal infections because some microorganisms can be found as a normal flora member and cause symptomatic disorder only in specific conditions. In addition, prevalence for vaginal infections is affected by method of diagnosis that can vary widely, from symptomatic questioning to cultures or molecular tests. In general, it is estimated that most of females will be involved with vaginal infection whilst their lifetime that is characterized by malodorous vaginal discharge, itching and burning (1).

Normal Vaginal Discharge

Vaginal discharge is the cardinal symptom for vaginal infections but may be difficult to differentiate the normal discharges from abnormal ones due to hormonal status of women. In reproductive ages, 1 to 4 mL of white or transparent, mostly odorless and not accompanied by itching, burning, irritation, erythema, local erosion and cervical or vaginal friability vaginal discharge is accepted as normal.

Vaginal nonkeratinized stratified squamous epithelium becomes rich in glycogen by effect of estrogen. Lactobacillus microbiom of vagina uses glycogen as a source that is shed from epithelium in order to provide acidic environment in the vagina that protects women from sexually transmitted diseases and opportunistic infections (2). In the premenarche and postmenopause, sparse amount of estrogen results in elevated vaginal pH (3). Also, vaginal normal discharge may become more apperent at pregnancy, ovulation and usage of estrogen-progestin contraceptives. Diet, sexual activity, medications and stress may impact on the volume and character of vaginal discharge.

Etiology

The most common causes of vaginal infections are vulvovaginal candidiasis, bacterial vaginosis and trichomoniasis (4,5). These infections are responsible for over 90 percent of infectious vaginitis (6). Also, cervicitis originating from sexually transmitted infections such as gonorrhea, chlamydia and mycoplasma may be found as nonspecific agents.

Common causes of vaginal infections will be discussed seperately below.

Candida Vulvovaginitis

Vulvovaginal candidiasis is one of the most common infections of vagina causing pruritus and leucorrhoea. It is described as vulvovaginal inflammation in the presence of *Candida* species. Since *Candida sp.* are part of normal vaginal flora of 25 percent of women (7), identifying the presence of *Candida* is not adequate to diagnose candida vulvovaginitis without symptoms.

Epidemiology:

Vulvovaginal candidiasis is the second most common cause of vaginitis symptoms and constitutes one-third of vaginitis cases (8). The exact prevalence is difficult to determine because *Candida sp.* not accompanied by inflammation can be identified in vaginal lumen and endocervical canal in 10 to 20 percent of healthy individuals in the reproductive age, 6 to 7 percent of menopausal women and 3 to 6 percent of prepubertal girls (9,10). The wide use of unprescribed antifungal drugs, diagnostic dependence on symptoms but not microscopic examination and cultures without clinical correlation can be listed as other causes for difficulty of epidemiologic studies. In a survey study, 55 percent of female university students reported at least one health care provider-diagnosed vulvovaginal candidiasis by age 25 (11) and 9 percent of women reported having four or more infections within 12 months, described as recurrent vulvovaginal candidiasis (12).

Microbiology and Pathogenesis

In 80 to 92 percent of vulvovaginal candidiasis cases in the United States, *Candida albicans* is identified and *Candida glabrata* is reported as the second most common cause for candidiasis (13). All *Candida* species cause similar symptoms but severity progresses milder in *C. glabrata* and *C. parapsilosis*. Access to vagina via migration from rectum across the perianal area is suspected since gastrointestinal tract and vagina cultures showing similar *Candida* species (14). Unlike bacterial vaginosis, vulvovaginal candidiasis is not associated with reduction of vaginal lactobacilli (15). Overgrowth of pathogen and penetration of epithelial cells is described as pathogenesis and severity of disease is associated with host inflammatory response and yeast virulence factors (16).

Risk Factors

Diabetes mellitus itself is a known risk factor for vulvovaginal candidiasis (17). Particularly nonalbicans *Candida* species have been identified pathogen among patients with diabetes mellitus type 2 (18). In the other hand, treatment of diabetes mellitus with sodium glucose cotransporter 2 inhibitors (such as canagliflozin, dapagliflozin, and empagliflozin) may increase the risk of colonization and inflammation of *Candida* species (19).

Usage of broad-spectrum antibiotics increases the risk of candida vulvovaginitis caused by decrease of normal vaginal microbiom (20). Besides, administration of oral or vaginal *Lactobacillus* during and after for four days after antibiotic use doesn't decrease the risk for development of postantibiotic vulvovaginitis (21).

Candida vulvovaginitis seems to be more frequent in circumstances progressing with elevated estrogen levels (For example: pregnancy and postmenopausal estrogen replacement therapy).

Immunosuppressed patients such as undergoing glucocorticoid or other immunosuppressive therapies or human with HIV infections are involved with increased prevalence of candidal infections (22).

Genetic studies have compared women suffering recurrent vulvovaginal candidiasis with control groups and identified polymorphisms in the *SIGLEC15* gene (23), *TLR2* gene and mannose-binding lectin genes (24), which are not modifiable risk factors but future gene therapy researches may decrease recurrence of vulvovaginal candidiasis.

Diet, sexual behavior, intrauterine or vaginal contraceptive devices and combined oral contraceptives are other known risk factors but studies have not reported consistent result about these risk factors, yet.

Symptoms and Physical Examination

Vaginal pruritus is the major symptom of vulvovaginal candidiasis (13). Vulvar burning and soreness are also common. Vulvar irritation may be present due to candidiasis itself or itching. Symptoms often worsen during the week before menses (25). The severity of symptoms varies from mild to severe except *C. glabrata* and *C. parapsilosis* infections, which is featured with minimal clinical findings or mild symptoms.

Erythema of vulva and vagina can be seen in physical examination. Vulvovaginal edema may be present. There can be minimal or no discharge. If present, vaginal discharge is classically white, thick and adherent to vaginal walls. It's been described as curd-like or cottage cheese-like, clumpy discharge. The malodorous discharge may be absent or minimal. The cervix usually appears to be normal.

Diagnosis

Diagnosis can be assessed via clinical findings, office evaluation, culture and nucleic acid tests. The office evaluation comprises of swabbing vaginal wall and discharge, assessing the vaginal pH and performing microscopy. The vaginal pH usually is normal but may be elevated with presence of concomitant bacterial vaginosis. In wet mount of discharge with addition of 10 percent potassium hydroxide, cellular components will be destroyed and microscopic evaluation will identify budding yeast, pseudohyphae and hyphae (Figure 1) (26). A surfactant, potassium hydroxyde and blue dye mixture known as Swartz-Lamkins fungal stain may make *Candida* structures easier to identify (27). Nevertheless, microscopy may be negative in 50 percent of cases with culture confirmed candidiasis (16).



Figure 1. Pseudohyphae of *Candida albicans* on wet mount vaginal discharge microscopy

The vaginal culture may be indicated in patients who are demonstrating clinical features of vulvovaginal candidiasis but no pathogens are present on microscopy, in order to avoid empiric

and unindicated therapies. The culture swab should be rubbed against vaginal sidewalls and inoculated onto Sabouraud agar or Microstix-Candida medium which performs equally well against first choice (13).

Polymerase chain reaction and other molecular tests may be considered in resistant and recurrent cases or in studies but clinicians should have regard to high costs and unnecessary of molecular tests in cases with highly specific clinical features. Also Pap smear may be positive in 25 percent of cases but it's an insensitive testing due to sample origin is cervix, which is not affected by *Candida* vaginitis.

Self-diagnosis is a common situation in vulvovaginal candidiasis and should be discouraged by health care providers. Women with prior infection usually make correct self-diagnosis but this will increase the unprescribed medication uses and may result with unnoticed and more serious gynecologic disorders.

Treatment

Vulvovaginal candidiasis treatment strategy should be set for relief of symptoms since 10 to 20 percent of reproductive aged females have *Candida sp.* as a part of vaginal flora, those do not require any therapy.

Treatment's severity is determined according to candidiasis, whether it's uncomplicated or complicated. Uncomplicated infection criterias are sporadic episodes (≤ 3 episodes/ year), mild to moderate symptoms, probable infection with *C.albicans*, healthy and non-pregnant patient, non-immunsuppressed individual. To tell that the infection is uncomplicated, all criterias must be present.

The initial treatment may be administered as fluconazole, 150 mg single dose. Therapeutic concentrations in vaginal secretions maintains for 72 hours after ingestion of single 150 mg tablet (28). When compared to topical antifungals, single-dose oral fluconazole regimen is less expensive treatment for candida vulvovaginitis.

Single dose topical regimens are also available in many countries for treatment of candidiasis. Topical agents seem to have less side effects, may lead to possible local burning and irritation, while oral treatment may cause gastrointestinal intolerance, headache, rash and transient liver function disorders. Also, oral treatment may take longer duration to relieve the symptoms. Oral or topical choice for therapy should be considered by cost, patient preference and contraindications. If the topical treatment has been chosen, not only vulvar but vaginal treatment would be crucial to eradicate candidal reservoir (29).

Ibrexafungerp is a single-day oral triterpenoid antifungal drug for use in uncomplicated vulvovaginal candidiasis. Ibrexafungerp shows its effect by inhibiting formation of fungal cell wall and resulting in death of *Candida* species. It's first in its class and may be indicated in patients who have allergic reactions or resistant infections to fluconazole or can not tolerate fluconazole therapy. Clinical trials have shown that ibrexafungerp results in high clinical cure rates compared to placebo, but comparison to azole treatment is yet to be recruited (30,31). It's been administered as two 150 mg tablets twice a day (i.e. two in the morning and two at night). CYP3A inhibitors may interact with ibrexafungerp, so patient who undergone CYP3A inhibitor treatment should reduce the dose to 150 mg twice in a day. Pregnancy and lactation data for humans are yet lacking.

In patients with severe symptoms or immunocompromise, fluconazole is suggested two or three (the number of doses depends on symptom severity) sequential 150 mg doses 72 hours apart.

C.glabrata has low vaginal virulence and infrequently causes vaginal symptoms. When suspected, adequate effort should be made to exclude other causes of symptoms. When *C.glabrata* is identified, vaginal 600 mg boric acid capsules once daily at night for two or three weeks may be administered. The success rates differ between 65 and 70 percent. For better results flucytosine

cream or 4% amphotericin B cream may be beneficial with combination of boric acid capsules (32). Neither boric acid capsules nor flucytosine or amphotericin B cream is available commercially and is made by compounding pharmacy. Boric acid capsules may be fatal if swallowed.

In pregnant women, clotrimazole topical agents should be used for one week because oral azole treatments may lead to miscarriage or birth defects, in first trimester, particularly.

Oteseconazole is another choice of azole antifungal drugs which is more potent than fluconazole in recurrent vulvovaginal candidiasis but has a long half-life (138 days) and contraindicated in women who have potential to get pregnant or lactating (33).

Trichomoniasis

Trichomoniasis is a genitourinary infection caused by protozan *Trichomonas vaginalis*. It's been known that trichomoniasis is the the most common nonviral sexually transmitted infection worldwide. It affects women more than men and is one of three common infectious agent that causing vaginal complaints. It's been estimated that trichomoniasis is affecting 3,7 million women and men in the United States annually and African-Americans are at risk greater four times compared to other races (34). World Helath Organization has estimated 156 million new trichomoniasis cases occurred in 2020 (35).

T.vaginalis is the flagellated protozan that is infecting squamous epithelium in the urogenital tract, especillay vagina, urethra and paraurethral glands (34). Cervix, bladder, Bartholin glands, prostate and even oral and rectal cavities are the other sites reported as involved.

Trichomoniasis is almost always sexually transmitted. Females acquire disease from both males and females, while males acquire disease from females but usually don't transmit disease to males. The incubation period is unknown but in-vitro studies have shown that incubation lasts for 4 to 28 days in approximately 50 percent of patients. Coexistence of trichomoniasis and bacterial vaginosis ise common, ranging from 20 to 80 percent. Screening for trichomoniasis is not recommended unless HIV infection is present.

Symptoms in female include a purulent, malodorous and thin vaginal discharge with burning, itching, dysuria, lower abdominal pain and dyspareunia. In physical examination, erythema of the vulva and vaginal mucosa may present. The classically describe yellow-green malodorous vaginal discharge occurs in 10 to 30 percent of women (36). Punctate hemorrhages may be shown on vaginal mucosa and cervix, named as strawberry cervix or colpitis macularis (Figure 2).

Untreated trichomoniasis may result in urethritis or cystitis, posthysterectomy cuff cellulitis and abscess, pelvic inflammatory disease, infertility, increased risk of acquiring HIV and HPV in non-pregnant women. In pregnant women, increased risks for premature rupture of membranes, preterm delivery and low birth weight infants have been reported.

Diagnostic evalution include wet mount microscopy and nucleic acid amplification test (NAAT). Clinicians should consider NAAT's longer turn-around time and higher cost and should prefer this test only in microscopic negative cases. In microscopy, jerky and spinning movement of flagellated *Trichomonas* may be seen in wet mount vaginal discharge preparates. Culture and cervical cytology is less accurate diagnostic tests and not recommended while trichomoniasis is highly suspected on clinical evalution (37).



Figure 2. Strawberry cervix or colpitis macularis seen in trichomoniasis

The treatment should be started with avoiding sexual activity until both male and female of couple are treated. Preferred therapy is metronidazole 500 mg orally twice daily for seven days. When it's not available, single dose of 2 g metronidazole is recommended but also associated with lower cure rates. Metronidazole vaginal gel is not recommended alone because doesn't treat all possible anatomic reservoirs.

Gonorrhea

Gonorrhea, the infection with the grma-negative coccus *Neisseria gonorrhoeae*, is a major cause of morbidities among sexually active women worldwide. Over 600,000 cases are reported in the United States annually (38). It's the major cause for urethritis in males and females. Thereafter the infection can result in pelvic inflammatory disease, infertility, ectopic pregnancy, chronic pelvic pain and disseminated gonococcal infection resulting with endocarditis and meningitis with severe morbidities. Gonococcal resistance for antibiotics is common.

The cervix is the most common site for mucosal infection in females. Up to 70 percent of cases may remain asymptomatic, thus the incubation period in females is less characterized than males. When symptoms are present, they manifest as vaginal itching and mucupurulent discharge. Some females may suffer intermenstrual bleeding and menorrhagia. Abdominal pain and dyspareunia may be present if the infection raises to upper genital tract. Urethritis may be presented with dysuria and urgency in posthysterectomy women, in particular. In association with pelvic inflammatory disease, perihepatitis (Fitz-Hugh-Curtis Syndrome) may exist and causes sharp pleuritic pain in the upper right quadrant of abdomen. Fitz-Hugh-Curtis syndrome is diagnosed usually incidental whilst surgery for any other reason. In pregnant women, gonococcal infections may result in chorioamnionitis, premature rupture of membranes, preterm birth, low birth weight and spontaneous abortions. Also, untreated *N.gonorrhoeae* infections may cause neonatal conjunctivitis in infants after delivery.

Treatment for uncomplicated cases of gonococcal infection is 500 mg ceftriaxone, administered intramuscular for individual weighed under 150 kg. Above 150 kg, the dose should revised as 1 g. If the presence of *Chlamydia trachomatis* can't be excluded, additive treatment with doxycycline (100 mg twice daily for seven days) is recommended due to high rates of coinfection.

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The Effects of Covid-19 on Musculoskeletal System and Pulmonary Rehabilitation

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Introduction

Coronavirus disease (COVID-19) is a disease which develops with respiratory tract symptoms; it was seen in Wuhan, China in December 2019 for the first time and was defined on January 13, 2020. The outbreak related with the disease first occurred in people in seafood and animal market in Wuhan region. Later on, it spread to other regions, especially Wuhan, to different provinces of China and then to many different countries (T.R. Ministry of Health COVID-19 Information Platform, 2020). Coronavirus (COVID-19), which gradually became a global health problem, affects the daily and social lives of individuals negatively. COVID-19, which initially occurred as a local health problem, became a global epidemic in a short time due to rapidly developing mobility and easy transmission (WHO, 2020). The disease is transmitted among people through the coughs and sneezing of sick individuals. Droplets scattering around as a result of the coughing and sneezing of the sick people and taking hands to eyes, face, nose or mouth without cleaning them after touching surfaces contaminated with respiratory particles are also the causes of virus transmission. Touching the face with contaminated hands increases the contagiousness of COVID-19 (T.R. Ministry of Health COVID-19 Information Platform, 2020). Individuals who have the disease have mild, moderate or serious symptoms. The disease, which usually targets upper respiratory tracts, also brings along musculoskeletal system diseases during and after the disease period. The disease has many side effects. These effects influence individuals and the society severely in health, social and economic areas. COVID-19 pandemic has affected all areas of life and also brought negative consequences in the world and in the field of health (Senol et al., 2021 ; TR. Ministry of National Education, 2020).

What are COVID-19 risk factors?

Not all individuals infected with COVID-19 are affected by the disease to the same degree. Age groups, profession and chronic diseases affect the disease, the course of the disease and its symptoms. 80% of individuals with the disease survive the disease with mild symptoms. 20% of individuals with the disease are treated in health institutions. COVID-19 has more effects on individuals aged 60 and older (T.R. Ministry of Health COVID-19 Information Platform, 2020).

Individuals most affected by the disease and age groups- Individuals older than 60 years of age: Although COVID-19 has affected all age groups, the elderly constitute the majority of individuals with the disease and deaths. According to data published by the United States of America in March 2020, it was reported that more than 80% of the disease-related mortality rate belonged to individuals over the age of 65 (Lee et al., 2020; Aytepe & Efe, 2021).

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Individuals with serious chronic medical conditions: In individuals with chronic disease, the process of surviving the disease and severity of the disease was more difficult than other individuals. Studies conducted showed that the most common comorbidities among chronic diseases were high blood pressure, diabetes, congestive heart failure, asthma, COPD (chronic obstructive pulmonary disease), inflammatory rheumatic disease and coronary artery disease (Günel et al., 2020).

Healthcare professionals: It was inevitable for healthcare professionals, who are at the forefront of the fight against the pandemic, to contact the virus directly. As of April 8, 2020, 22.073 healthcare professionals diagnosed with COVID-19 from 52 different countries were reported to World Health Organization (Pala & Metintaş, 2020).

What are the signs and symptoms of COVID-19?

Common symptoms of COVID-19 are high fever, fatigue and weakness, cough, loss of sense of taste or smell, muscle, joint and back ache. Less common symptoms are headache, diarrhoea, sore throat, skin exfoliation, redness and irritation of the eyes. Severe symptoms are respiratory distress or shortness of breath, confusion, disorientation in movement or speech and chest pain.

What are the post COVID-19 musculoskeletal system complaints?

Low back, back, shoulder and neck pain are common post COVID-19 musculoskeletal system complaints. A large number of studies conducted support these symptoms. The regions with more intense pain in the musculoskeletal system have been found as neck, pain and low back region (Gayretli et al., 2021). Studies conducted have shown that the region where pain occurs most in actively working people is the formations on the spine as the waist, neck and back. In one study, it was stated that waist, back and neck regions had the highest rate of pain during the last year, highest rate of work disability during the last year and the highest rate of pain during the last week (Jang et al., 2014; Briggs et al., 2009). A study by Skovlund et al. (2020) researched musculoskeletal pain in working population and stated that pain occurred mostly on waist and neck regions (Skovlund et al, 2020).

Post COVID-19 Management of Pain in Musculoskeletal System

The main effect in COVID-19 is lung involvement. However, studies conducted have stated that some other organs are also targeted because they have some common tissues with the lungs (ACE 2 receptors). These receptors targeted by the virus can be found in intestines, small vessel endothelium, smooth and striated muscle and synovial tissue. Therefore, when the existing cases are examined, it is possible to encounter not only cough, dyspnoea or high fever, but also symptoms of fatigue, myalgia and arthralgia. The prognosis is no doubt more severe when the patient's motility loss, malnutrition and respiratory problems are added to this picture.

While musculoskeletal system pain is one of the most common acute findings of COVID-19 infection, chronic complications related with moderate and severe COVID-19 have also been reported. In addition to prolonged immobilization and intensive care units processes, pro-inflammatory, cytokines and chemokines and direct damage of SARS-Cov2 to the tissue have been held responsible for the destruction of skeletal muscle, bone, joint, nerve and soft tissues (Huang, 2020). In studies conducted, while the complaint of weakness was the complaint that continued most frequently with a rate of 58% in post COVID-19 patients, joint complaints were found to occur in 19% of the patients (Lopez-Leon, 2021). In addition to this, it was reported that the frequency of muscle weakness or fatigue was 63% in post COVID-19 sixth month (Huang, 2021). Some of the pain felt by patients is related to the pathophysiology. However, the problems in posture and bodily awareness that occur due to lack of movement and spending more time with

technological and electronic devices during the isolation period should not be overlooked. Clinicians should also accept the importance of describing pain phenotypes correctly. For example, it has been suggested that infections may affect peripheral nerve system or central nerve system directly or induce post-viral immune syndrome supporting the development of neuropathic pain. Patients with neuropathic pain usually have negative results when compared with patients who have nociceptive pain. For this reason, evaluating pain phenotypes in patients with permanent COVID-19 symptoms provide important information for healthcare professionals. In addition, the treatment protocol to be applied guides the choice of methods and exercises that will provide holistic healing as well as the muscle and bone structures that are symptomatic (Karateev, 2021). Many problems continue in patients after COVID-19 infection. Patients may have respiratory symptoms such as fatigue, weakness in skeletal and respiratory muscles and atrophy, decreased exercise capacity, fatigue, pain, limitation of functions, dyspnoea during low intensity exercise or daily living activities; psychosocial problems such as anxiety, depression and post-traumatic stress disorder; and prolonged problems such as cognitive disorders, speech and swallowing disorders, deterioration in daily living activities and decreased quality of life. Therefore, COVID-19 patients are candidates for pulmonary rehabilitation (PR) (Zhao et al., 2020; Yang & Yang, 2020).

Methods used in pulmonary rehabilitation for post-COVID patients

Physiotherapy agents

1. Tens (Transcutaneous Electrical Nerve Stimulation): High or low frequency tens is a physiotherapy agent that has been found to be very effective especially in waist and back pain in studies conducted. Tens can be preferred especially for short-term immediate relief of pain (Oral & Ketenci, 2013).

2. Therapeutic ultrasound: These are devices producing high frequency sound waves. Sound waves can disperse in solid and liquid space. Heat energy is released by reflection in the tissues through which these waves pass. Sound waves have the ability to disperse in solid and liquid space. Heat energy is released by reflection in the tissues through which these waves pass. Micro-massage and warming effect of ultrasound is used in the treatment of pain. The skin is nourished and revitalized by stimulating and working in the capillaries. Ultrasound may be preferred for the recovery of soft tissue disorders and relief of arthralgia symptoms. No treatments are made on bone tissue (Oral & Ketenci, 2013).

3. Hot application: It is the warming of skin and subcutaneous tissue with the help of special fabric bags that do not leak water. A more superficial heat effect is released when compared to ultrasound. The veins expand and circulation relaxes with the effect of increasing heat. It is not preferred in acute inflammation and rheumatological cases and also in cases metastasis may occur (Özdinçler, 2016)

Therapeutic Exercises

Therapeutic exercise is a physical therapy technique used to improve or maintain an individual's physical condition through resistance, endurance, flexibility and balance training. The intensity, volume, progression and type of exercises should be personalized based on the physical condition and tolerance while performing the exercise. When prescribing exercise to patients, exercise type, intensity and frequency suitable for the result of the evaluation should be determined. It has been stated that exercise therapy will be beneficial in the recovery of muscle, bone, joint and connective tissue during the COVID-19 process (Öztürk & Kaya, 2020).

1a. Core stability and balance exercises: The ability of muscles to provide the functional stability of joints by creating synchronized activity and controlled movement is called neuromuscular control. The part of the body known as “core”, or the lumbopelvic hip component,

consists of both passive and active structures of the hip and trunk. Stabilization is mainly achieved by co-contraction of the muscles. Core stabilization is very important in maintaining the balance of spine structure. A healthy posture supports the position and functions of the tissue and organs. It also provides the adequacy of joint range of motion in the neck, upper and lower extremities (Yüksel et al., 2019).

1b. Mobilization: According to clinical studies conducted, in and out-of-bed activities performed according to the respiratory and muscular capacity of the individual reduce pain significantly. Transition from lying in bed to sitting position, sitting and short distance walking can be given as examples to these activities (Kırmızı, 2020). Early physiotherapy applications including gradual mobilization and exercise therapy have an important place in reducing complications associated with immobilization, returning to daily life and increasing physical activity tolerance in COVID-19 patients (Kırmızı, 2020).

1c. Joint range of motion exercises: They are the movements performed by the muscle and joint by using the physiological limits of the joint range of motion. They can be performed passively or actively (Yüksel, 2019).

1d. Aerobic exercises: Exercises such as walking, running, swimming and cycling are aerobic exercises; they work the major muscles and have effects such as increasing aerobic capacity, decreasing insulin resistance and inflammation and losing weight. These effects have been found to be effective in decreasing pain intensity in patients with musculoskeletal system pain (Meng, 2015). In patients with continued oxygen support, low intensity aerobic exercises such as walking (<3 MET) are started with close monitoring of blood pressure, SaO₂, heart rate monitorization and newly developed cardiorespiratory symptoms. Duration and intensity of exercise are gradually increased according to the patient's condition by starting 3-4 times a week and for 10-15 minutes (Barker-Davies, 2020).

1e. Strengthening exercises: 1 repetitive maximum (RM) 30-40% mild and moderate intensity 1-3 sets of exercise, 8-15 repetitions of each set with 2 minutes of rest in-between sets should be performed 2 times a week by starting with 3-4 muscle groups in very weak patients. Weight should be lifted for 3-6 seconds without causing pain and discomfort. Valsalva manoeuvre should be created by exhaling while lifting (Yüksel, 2019). Intensity of exercise and number of repetitions are gradually increased according to the tolerance and condition of the patient. While returning to exercise from mild/moderate severity COVID-19, 1 week of low level stretching and mild muscle strengthening activity should be tried before targeted cardiovascular sessions (Barker-Davies, 2020).

1f. Diaphragm exercises: In patients who have had COVID-19, the aim of respiratory exercises is to regulate the respiratory cycle, improve dyspnoea symptoms, regulate respiratory movements, increase oxygenation, decrease secretion in airway, prevent dysfunction, decrease anxiety and depression, provide and improve relaxation, maintain functions and increase quality of life (Arzani, 2020; Zhao, 2020). The objective of PR is not only to improve the physical condition of individuals. It also enables individuals to return more quickly to their environment and the society (Yang & Yang 2020).

With the correct use of diaphragm muscle in individuals who have had COVID-19, the capacity and efficiency of respiration increases, organ mobility is ensured, lymphatic system is activated, spine mobilization is increased with chest mobility and the rate of healing increases in tissues and organs (Yang & Yang 2020).

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X-Ray Effects on Chondrocyte Volume from Bovine Articular Cartilage

Ekrem ÇİÇEK¹

Introduction

The amount of ionizing radiation that a person is exposed to has sharply risen over the last decades (Leuraud & et al., 2015). Medical radiation exposure is the largest part of the total amount of radiation. The average yearly medical exposure dose had increased almost 600% (0.5 mGy in 1982 and 30 mGy in 2006) in the United States (Mettler & et al., 2008). The largest parts are coming from nuclear medicine applications and computed tomography scanning. Occupational sources of radiation exposure are also important (Leuraud & et al., 2015; Mettler & et al., 2008).

The effects of X-ray radiation on the mechanical properties of articular cartilage was studied previously. It was found that ionizing radiation caused adverse effects on the function of articular cartilage. The Young's Moduli for the controls were significantly higher than the ionizing radiation exposed samples (Cash & Dean, 2019; Lindburg, Willey & Dean, 2013). The same effects of ionizing radiation found by other studies (Cicek, 2016; Kuzu & Cicek, 2019). Ionizing radiation increased cellular apoptosis both in vivo and in vitro studies (Rocha & et al., 2016).

The effects of ionizing radiation on articular cartilage are characterized as negative effects, no effects and positive effects in the literature (Cash & Dean 2019). Previous studies on cartilage response to ionizing radiation are inconsistent and radiation effects on articular cartilage is unclear (Hugenberg, Myers & Brandt, 1989; Willey & et al., 2013).

Articular cartilage is the smooth connective tissue and is 2 to 4 mm thick (Sophia, Bedi & Rodeo, 2009). The chondrocytes are the only cells in articular cartilage. Chondrocyte shapes are depth-dependent in articular cartilage. There are flat chondrocytes, spherical chondrocytes and elongated chondrocytes in the superficial, middle and radial zone, respectively (Sophia, Bedi & Rodeo, 2009). The injury reaction of the tissue is inefficient since articular cartilage is an avascular tissue (Willey & et al., 2013).

It is important to know ionizing radiation effects to balance medical applications against the risks. The aim of this study was to investigate the effects of ionizing radiation on chondrocyte volume from bovine articular cartilage. The size of the chondrocyte was measured from confocal image of articular cartilage.

Materials and Methods

Explant harvest

Mature bovine knees with joint capsules were obtained from a local abattoir (Bud's Custom Meats, Riverside, IA, USA). Osteochondral explants (2.5 cm x 2.5 cm) were carefully excised from the lateral tibial plateaus of the joints under sterile conditions. Then, they were placed immediately in Hanks balanced salt solution (HBSS). HBSS was supplemented with penicillin/streptomycin and amphotericin B. Explants were incubated in culture medium during the whole study to maintain tissue viability (Ramakrishnan & et al., 2011; Cicek & Arıkanoglu, 2014; Goodwin & et al., 2010; Martin & et al., 2012).

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X-ray Irradiation

The explants were exposed with X-ray 150 keV, 15 mA. The distance between X-ray source and samples was 115 cm. The total exposed dose was 100 mGy. Imaging process started 3 hours after the irradiation.

Confocal imaging

The explants were stained with dihydroethidium (DHE, Invitrogen) and calcein AM (Invitrogen) for 30 min. The stained explants were scanned on a BioRad 1024 Confocal Microscope equipped with a Krypton/Argon laser (Biorad Laboratories Inc., Hercules, CA). Images of the osteochondral explants were performed with a 20× water immersion lens. The random sites close to the radial center were imaged from the surface to a depth of ~200 µm in 20µm intervals. 20× image stacks were acquired from randomly selected imaging sites. 20× image stacks were utilized for chondrocyte shapes analysis. More information regarding imaging conditions was described previously (Ramakrishnan & et al., 2011; Cicek & Arikanoglu, 2014; Goodwin & et al., 2010; Martin & et al., 2012).

Image Analysis

The ImageJ version 1.53 was used to analyze live cells from confocal image stacks. We measured the surface area of the chondrocyte, perimeter, long (major) and short axis (minor) inside the cell, height of cell, cell angle, circularity of the chondrocyte and aspect ratio. The size (area) of the chondrocyte was selected between 20 to 80 µm². The circularity of the chondrocyte was among 0.01-1.00. ImageJ version 1.53 for Windows was obtained from the NIH website (ImageJ, Rasband, W.S., U. S. National Institutes of Health, Bethesda, MD, <http://rsb.info.nih.gov/ij/>).

Results

Table 1 presents the chondrocytes parameters for the control group.

Table 2 shows the chondrocytes parameters for the X-ray exposed group.

The p values show the significance between the control group and X-ray exposed group. The volume ($p = 0.001$), area ($p < 0.001$), perimeter ($p < 0.001$), major axis ($p < 0.001$) and minor axis ($p = 0.006$) were significant. The angle of chondrocytes ($p = 0.820$), circularity ($p = 0.139$) and aspect ratio ($p = 0.343$) were insignificant.

Table 1. The chondrocytes parameters for the control group.

Depth (µm)	Number of cells	For each depth the mean values and standard errors of chondrocytes							
		Volume (µm ³)	Area (µm ²)	Perimeter (µm)	Major axis	Minor axis	Angle	Circularity	Aspect ratio
0	30	1496.92 ± 46.95	86.74 ± 2.51	37.29 ± 0.63	13.32 ± 0.36	8.38 ± 0.23	88.19 ± 8.96	0.79 ± 0.02	1.65 ± 0.08
-40	30	1205.31 ± 67.68	80.79 ± 4.32	35.50 ± 1.13	12.91 ± 0.55	7.96 ± 0.29	100.86 ± 9.39	0.80 ± 0.02	1.69 ± 0.11
-80	30	1247.07 ± 60.74	84.07 ± 4.04	35.66 ± 0.97	12.99 ± 0.42	8.17 ± 0.24	77.74 ± 7.13	0.82 ± 0.01	1.62 ± 0.06
-120	30	1130.03 ± 51.16	75.17 ± 3.31	33.75 ± 1.01	12.27 ± 0.51	7.84 ± 0.23	80.11 ± 9.00	0.83 ± 0.02	1.61 ± 0.09
-160	30	1168.24 ± 57.58	78.00 ± 3.89	34.57 ± 1.27	12.16 ± 0.53	8.16 ± 0.22	90.38 ± 7.88	0.83 ± 0.02	1.53 ± 0.09
-200	30	1221.57 ± 73.98	73.62 ± 4.10	34.86 ± 1.38	12.66 ± 0.59	7.39 ± 0.23	86.63 ± 8.26	0.77 ± 0.02	1.76 ± 0.10

Table 2. The chondrocytes parameters for the X-ray exposed group.

Depth (µm)	Number of cells	For each depth the mean values and standard errors of chondrocytes							
		Volume (µm ³)	Area (µm ²)	Perimeter (µm)	Major axis	Minor axis	Angle	Circularity	Aspect ratio
		p = 0.001	p < 0.001	p < 0.001	p < 0.001	p = 0.006	p = 0.820	p = 0.139	p = 0.343
0	30	851.75 ± 49.59	56.93 ± 3.31	29.03 ± 0.97	10.50 ± 0.40	6.79 ± 0.22	103.10 ± 9.55	0.84 ± 0.01	1.57 ± 0.06
-40	30	980.21 ± 56.31	65.64 ± 3.80	31.18 ± 0.96	11.13 ± 0.41	7.40 ± 0.22	105.58 ± 9.14	0.83 ± 0.01	1.52 ± 0.15
-80	30	996.36 ± 51.96	66.92 ± 3.43	32.30 ± 0.97	11.93 ± 0.44	7.21 ± 0.28	94.80 ± 11.64	0.81 ± 0.01	1.70 ± 0.08
-120	30	938.98 ± 43.59	63.48 ± 2.87	31.17 ± 0.86	11.36 ± 0.39	7.10 ± 0.22	78.82 ± 10.87	0.82 ± 0.02	1.65 ± 0.08
-160	30	999.35 ± 40.22	66.93 ± 2.69	31.94 ± 0.80	11.63 ± 0.42	7.39 ± 0.23	79.22 ± 9.97	0.83 ± 0.02	1.64 ± 0.10
-200	30	1055.16 ± 51.15	70.84 ± 3.41	32.72 ± 0.89	11.47 ± 0.35	7.82 ± 0.26	71.76 ± 9.86	0.82 ± 0.01	1.51 ± 0.07

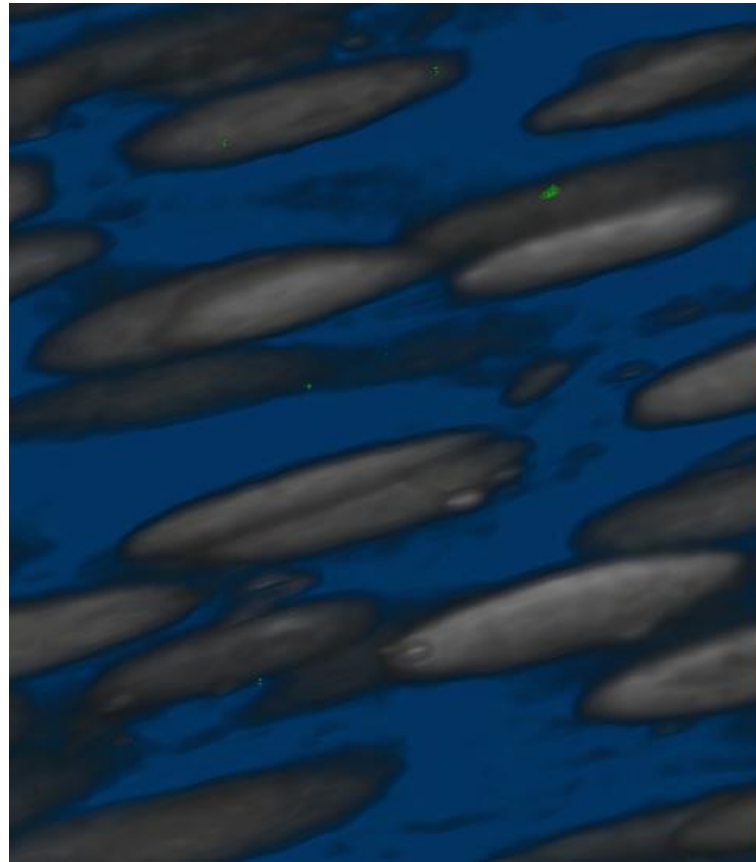


Figure 1. Confocal image from bovine articular cartilage

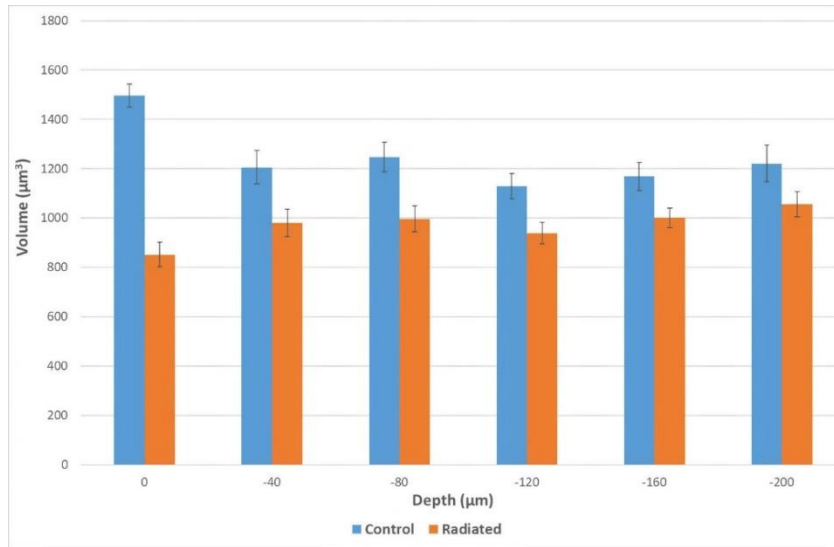


Figure 2. The depth-dependent volume of chondrocyte for control and irradiated groups

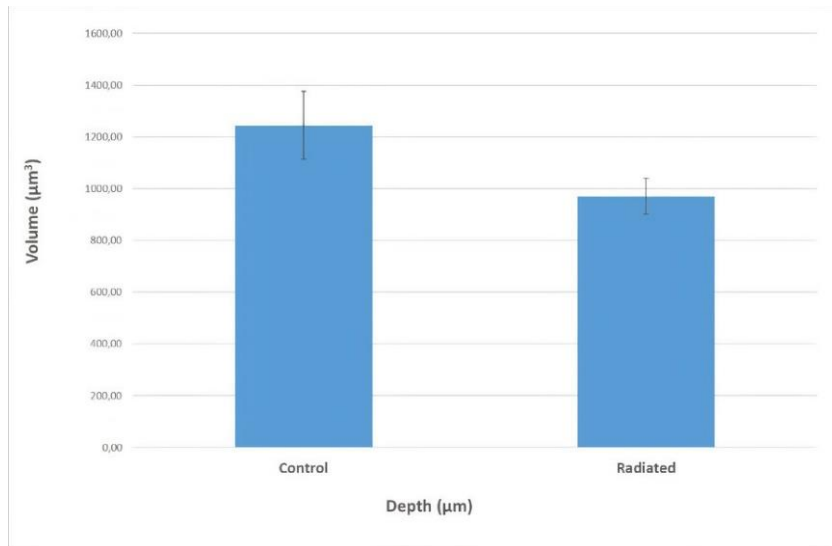


Figure 3. The mean value of chondrocytes volume for control and irradiated groups

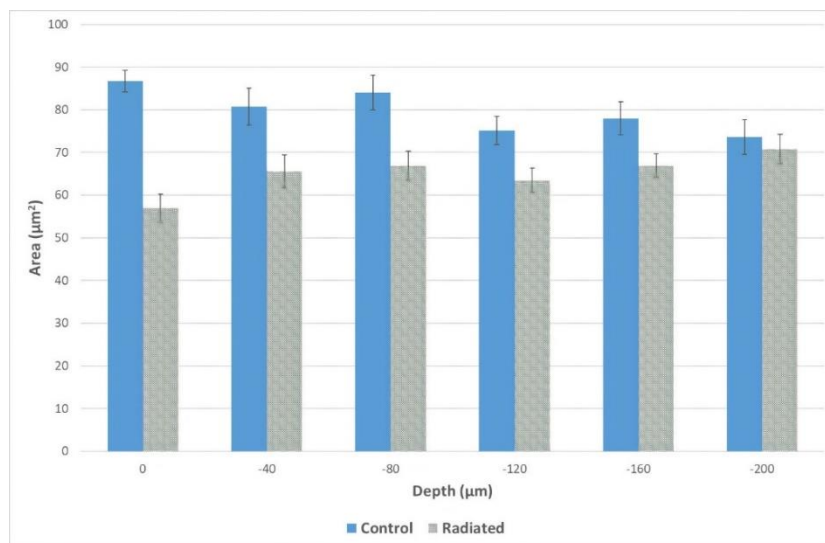


Figure 4. The depth-dependent area of chondrocyte for control and irradiated groups

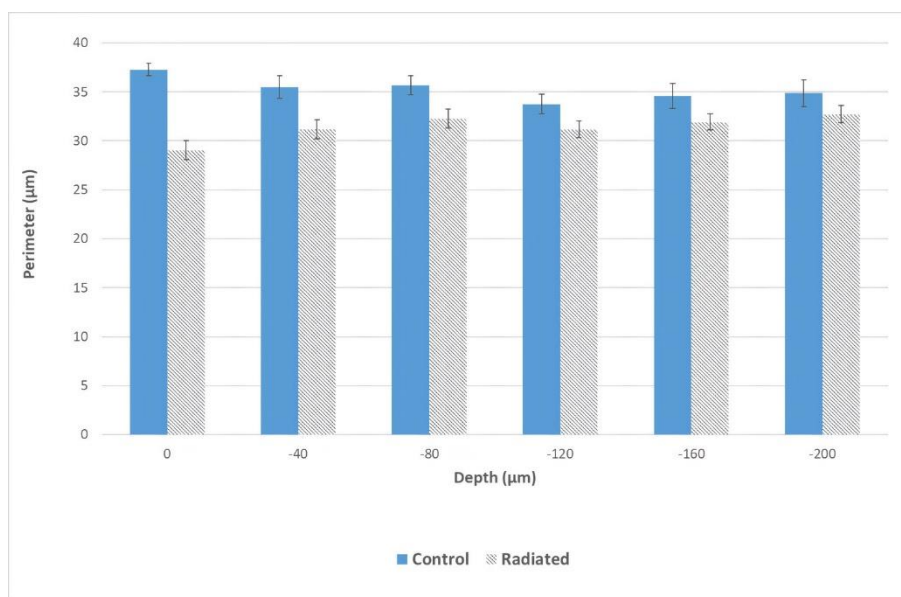


Figure 5. The depth-dependent perimeter of chondrocyte for control and irradiated groups

Figure 1. shows confocal image from bovine articular cartilage. The volume of the chondrocyte can be seen in this image. Figure 2 demonstrates the depth-dependent volume of chondrocyte for control and irradiated bovine articular cartilage. The chondrocyte volume in the control group is higher than the X-ray exposed group for all depth of articular cartilage. The lowest and highest mean volume of chondrocytes were 852 μm^3 and 1055 μm^3 , respectively for the exposed group.

Figure 3 presents the mean value of chondrocytes volume for control and irradiated bovine articular cartilage. The mean value of chondrocyte volume in the control group was 1245 μm^3 , while it was 970 μm^3 in the exposed group. The mean value of chondrocyte volume in the control group was significantly higher than the X-ray exposed group. The mean value of chondrocyte volume was significant between the groups ($p=0.001$).

Figure 4 shows the depth-dependent area of chondrocyte for control and irradiated bovine articular cartilage. The chondrocyte area in the control group is higher than the X-ray exposed group for all depth of tissue.

Figure 5 demonstrates the depth-dependent perimeter of chondrocyte for control and irradiated bovine articular cartilage. The chondrocyte perimeter in the control group is higher than the X-ray exposed group for all depth of cartilage.

Statistical analyses were performed using the Minitab 19 software. One-way analysis of variance (ANOVA) was used for statistical analysis. A p -value less than 0.05 was considered significant. The depth-dependent measured chondrocyte volume was demonstrated for control and radiated groups. The volume of chondrocytes was decreased in the exposed group. The results were significant ($p=0.001$).

Discussion

Reduction in matrix metabolism is displayed as the reason of cartilage weakening at articular surfaces in response to irradiation which, due to permanent deterioration of mechanical properties, may contribute to overall joint erosion (Willey & et al., 2013), joint failure (Kolár, Vrabec & Chyba, 1967) and osteoarthritis (Goldring & Marcu, 2009).

It has been shown to induce senescence among young rabbit articular chondrocytes early after radiation exposure (Hong & et al., 2010). It has been reported that radiation exposure of rabbit knee with a total dose of 5 Gy does not affect cartilage mechanical properties, although chondrocyte clustering changes were seen (Öncan & et al., 2013).

Since types and exposure amounts are varying, ionizing radiation effects on articular cartilage is not clear. Degradation of cartilage following exposure may be dependent on the radiation model and radiation dose (Hugenberg, Myers & Brandt, 1989; Jikko & et al., 1996). Ionizing radiation (2 Gy and 10 Gy doses) induced active degradation of matrix and reduced proteoglycan synthesis in human and pig cartilage and chondrocytes. However, the nature of the degradation and impaired production of proteoglycans following ionizing radiation exposure is not clear (Willey & et al., 2013).

A major pathway of active degradation of articular cartilage after radiation exposure may be acute loss of glycosaminoglycans (Cornelissen, Thierens & De Ridder, 1996). The glycosaminoglycan content a week after the irradiation in a study was 55% lower in irradiated explants than controls, in which recovery from the initial loss is not likely (Rojas & et al., 2014). It was shown that 2 Gy X-ray exposure reduces stiffness in a 3D model of porcine articular cartilage. This reduction is related with a large release of glycosaminoglycans from irradiated explant 24 h after exposure (Saintigny & et al., 2005). It is believed that glycosaminoglycan loss is not the only factor affecting the modulus after radiation exposure, although other causes remain undefined. It was not possible to determine if changes in cross-linking or other structural damage are contributing to the reduction in modulus after radiation exposure (Lindburg, Willey & Dean, 2013; Willey & et al., 2013; Hong & et al., 2010).

It was reported that X-ray exposure with 20 Gy decreased type II collagen and glycosaminoglycan production (Cornelissen, Thierens & De Ridder, 1993). They also showed that 50 Gy X-ray dose reduces dramatically extracellular matrix components in 14 days in mature cartilage (Cornelissen, Thierens & De Ridder, 1993a, 1993b). It was shown that type II collagen reduction after X-ray exposure is linked with induction of type I collagen, especially into pericellular spaces of chondrocytes (Muhonen & et al., 2006). It was proposed that radiation-induced changes in oxygen concentrations might increase in type II collagen. Furthermore, it has been suggested that reactive oxygen species (ROS) suppress proteoglycan synthesis. Radiation effect on proteoglycan synthesis might reflect effects of ROS, since free radicals suppress proteoglycan synthesis (Bates, Johnson & Lowther, 1985a, 1985b).

These dramatic failures of fibril organization cause a loss of mechanical properties in mature cartilage (Muhonen & et al., 2006). Moreover, it causes a functional decline of cartilage health, in patients treated with ionizing radiation (Cornelissen, Thierens & De Ridder, 1996; Rojas & et al., 2014). Constant radiation caused changes in extracellular matrix metabolism could deteriorate cartilage mechanical properties after exposure, by extracellular matrix degradation (Saintigny & et al., 2005).

It was reported that lower doses of ionizing radiation may cause a significant decrease in the compressive stiffness of articular cartilage at multiple length scales. Young's modulus reduction in mechanical properties following radiation exposure suggests that alterations are occurring in the matrix. This reduction in Young's modulus after radiation exposure is not solely due to the observed loss of proteoglycans, which are only estimated to be responsible for about 50% of the compressive stiffness of cartilage (Buschmann & Grodzinsky, 1995).

In a previous study, application of X-ray at 4.8 mGy in compressive moduli of equine articular tissue experiments produced similar results with another study (Kuzu & Cicek, 2019), in the decreasing of stiffness (Cicek, 2016).

In a previous study (Bates, Johnson & Lowther, 1985b), in which the differential effect of C-ions versus X-rays was investigated in human articular cartilage, exposure to X-ray and C-ions were not able to induce neither necrosis nor apoptosis in either 2D or 3D culture. The previous studies are supporting the relative radiation resistance of chondrocytes, regardless of the radiation quality in rabbit articular chondrocytes (Hong & et al., 2010), porcine and human articular cartilage (Willey & et al., 2013). However, exposure to ionizing radiation might induce bone differentiation and senescence (Saintigny & et al., 2005), the latter is an important factor of potential cartilage attrition (Hamdi & et al., 2016).

Conclusions

Application and exposure of ionizing radiation is increasing globally. Articular cartilage is the thin and white connective tissue. Chondrocytes are the cells in articular cartilage. The effects of ionizing radiation depend on type and amount of ionizing radiation. In this study, ionizing radiation effects on chondrocyte volume was investigated. We also investigated the changes of chondrocyte shape after ionizing radiation exposure. The results were reported according to the depth of chondrocyte since it displays zonal shape dependency. Further research is necessary to identify potential molecular targets for the functional deficits in cartilage post-irradiation. In conclusion, chondrocyte volume of bovine articular cartilage was found to reduce after exposure to ionizing radiation with X-ray. We demonstrated ionizing radiation (X-ray) effects on chondrocyte volume from confocal image of bovine articular cartilage.

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Rational Use of Medicaments

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Introduction

The definition of rational drug use was first discussed at a meeting held in Nairobi, Kenya in 1985. According to the definition made here, rational drug use; “It is all of the rules that require patients to take the drugs they need to use in accordance with their clinical needs, in an amount that meets their personal needs, in a sufficient time frame, at the lowest cost to themselves and the society” (Holloway & Dijk, 2011).

The use of drugs, which cannot be considered rational, is known as a type of habit that is difficult to correct. Although irrational drug use seems to be a serious problem in all countries, it is more common in underdeveloped or developing countries (Acar & Yeğinoğlu, 2005). In addition to insufficient education level, economic, sociocultural and regulatory mechanisms are important factors in irrational drug use (Kaya et al., 2015).

The issue of rational drug use plays an important role in the interaction of individuals' personal health and social and economic structures and being an effective element of the health care management system (Amin et al., 2011).

Various factors are effective on rational drug use. One of the most important of these is the level of health literacy. It has a direct relationship. These two elements are concepts that complement each other and one improves the other. Individuals with a high level of health literacy are also expected to have a high level of rational drug use (İncesu, 2017).

World Health Organization for rational drug use is guiding in the drug treatments that physicians will recommend to patients and in points such as drug prescribing. This guideline includes criteria such as being effective, safe, appropriate and economical when choosing drugs. These issues are very important in prescribing drugs to patients (Gülhan, 2013; Erden & Göçmez, 2004; Akıcı et al., 2002).

All behavior, except for the rational use of drugs, can be described as irrational drug behavior. Irrational drug use behavior emerges as a result of patients not getting enough benefit from drugs. This situation creates an extremely negative impact on the economies of countries (Ekenler & Koçoğlu, 2016).

Many processes are carried out to ensure the effective, efficient and functional delivery of health services. Due to the scarcity of resources, their efficient use and the implementation of policies aimed at saving have gained importance. Thus, with the increasing health needs, the importance of making rational decisions about using scarce resources more effectively increases (Ateş & Aba, 2019). Living healthy is the most basic right of every human being. For this reason, it is the most natural right of individuals to want to be treated when their health integrity is impaired (Sevim & Nal, 2021). Policies for rational drug use are important for the sustainability of this system.

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Rational Drug Use

The first step in rational drug use is the creation of the Primary Essential Medicines list, which was determined by the World Health Organization (WHO) in 1977 to assist in the preparation of national drug lists to be created in line with the needs of countries. Rational drug use, with its current definition; In a meeting organized by WHO in Kenya in 1985, it was stated as *“The set of rules that require patients to take drugs suitable for their clinical needs, in doses that meet their personal needs, for a sufficient period of time, at the lowest cost for themselves and the society”*. Following this first step, the International Network for Rational Drug Use (INRUD) was established in 1989 in order to carry out multicenter research projects from a single center and to support the rational use of drugs (WHO, 2002).

Ministry of Health, rational drug use; It defines it as the planning, execution and monitoring process that enables the use of drugs in an economical, effective and safe manner (TC Ministry of Health, General Coordination of Health Project, 1993). According to the definition of the World Health Organization in 2002, it is the planning process in a way that can provide the rational use of drugs according to the clinical results of the individuals, in the appropriate dose according to their individual characteristics, at the lowest cost for a sufficient period of time (WHO, 2002).

The primary goal in rational drug use is to prevent all the physiological, biological and psychological problems that may arise with the wrong use of drugs, and to try to minimize social and economic damage as much as possible (Ulusoy & Sumak, 2011). While one-third of the world's population struggles to purchase even essential drugs, the ease of access to drugs by some people without a prescription, and the increase in the amount and types of drugs with each passing day show the necessity of the concept of rational drug use (WHO, 2002).

According to the World Health Organization, a drug is defined as *“a substance or product that is used or intended to be used to modify or examine physiological systems or pathological conditions for the benefit of the recipient, and to provide mental or physical well-being in the user”* (WHO, 2002). Medicines are an essential component of health. In this context, it plays a vital role in healing patients and saving lives (Angamo, Wabe & Raju, 2011).

The use of drugs has an important place in the health services offered in terms of the benefits it will provide in the lives of people in the cure of many diseases and in the mortality and morbidity rates caused by many diseases, or vice versa, in the correct and sufficient consumption (Büyükturan & Büyükturan, 2017).

The drug should be used rationally, not randomly. In addition to the economic burden it reflects on individuals and society, unconscious drug use can even result in death. A drug is a product used to protect or regain health, rather than irrational consumption. For this reason, it is necessary to consume the drug consciously and in sufficient quantities (Akıcı et al., 2011; Kurt et al., 2020).

Medicines have an undeniable importance in protecting, maintaining and improving the health of individuals (Memişoğlu, 2016). If drugs and treatments are not managed correctly, they can cause many health problems and death, in addition to their protective and curative effects. Accordingly, the increase in medical costs makes the rational use of drugs important (Pinar, 2012).

all over the world and in Turkey is a concept whose importance is increasing day by day because the country's resources are not wasted. Compared to developed countries, the amount of drugs used in Turkey is stripped of its therapeutic dimension and takes the form of wasting more resources. In this context, the most important cause of waste in drug consumption is over-the-counter and unconscious consumption. Compared to other countries, the share allocated only to

pharmaceuticals in healthcare payments is 25-30% higher in Turkey (Demiroğlu, Polat & Doğan, 2017).

Research shows that more than half of the drugs in the world are prescribed by doctors, but only half of individuals use these drugs correctly in accordance with recommendations. Considering these rates, it is understood how important rational drug use education is (WHO, 2002; Holloway & Dijk, 2011).

Basic Principles of Rational Drug Use

The prescribing guide, organized by the World Health Organization, is a guide in the drug treatments that physicians will recommend to patients and in points such as drug prescribing. This guideline includes criteria such as being effective, safe, appropriate and economical when choosing drugs. Physicians need to create a list of drugs and treatment models that are compatible with the patient, taking into account the prescribing guidelines (Gülhan, 2013; Erden & Göçmez, 2004; Akıcı et al., 2002).

- Effectiveness: Pharmacological properties of drugs
- Being safe: Side effects due to drug use
- Eligibility: Side effects caused by drugs
- Being economical: Presentation price of drugs

Rational Drug Use Criteria

In order to choose an effective, safe, appropriate and economical drug in the principles of rational drug use, a set of criteria must be met in the planning, functioning and follow-up processes. Drug rationality is to ensure that evidence-based information produced by experimentation and observation is effectively included in the diagnosis and treatment process, and is accessible in terms of financing drugs and when necessary. In this context, certain criteria are needed. Criteria for rational drug use; rational choice of drugs, rationality of the pharmaceutical industry, the process of prescribing and responding to prescriptions, rational consumption of drugs, and the rationality of regulations on drug management (The Alphabet of Rational Drug Use, 1993, www.e-kutuphane.teb.org.tr).

Rationality of Pharmaceutical Logistics: It consists of purchasing, storage, inspection and distribution of drugs. Rational financing in the purchasing process of the drug, storage and storage in appropriate conditions, delivery to the point of need in a timely and sound manner, and ensuring that all these steps are properly supervised are among the suggestions regarding the rationality of drug logistics (Uslu & Akçadağ, 2012; The Alphabet of Rational Drug Use, 1993, www.e-kutuphane.teb.org.tr).

Rational Selection of Medicines: The physician should make a “rational” (rational) drug selection in accordance with the clinical condition of the patient, of the drugs whose safety and effectiveness have been tested and proven in line with the generally accepted parameters. If a rational choice cannot be made in this regard, ineffective treatments will emerge, leading to unnecessary drug consumption by patients (Sağır & Parlakpınar, 2014). It is recommended to consider standardized treatment protocols, epidemiological data obtained with mortality and morbidity numbers, and the distribution of patient burden in determining the priority drug list (The Alphabet of Rational Drug Use, 1993, www.e-kutuphane.teb.org.tr).

Rationality of the Prescribing Process: It is the stage where the physician participates in the rational drug use process. In this process, physicians are the party directly responsible. This stage should begin with the correct diagnosis (Yorulmaz, 2003). This process, which is a part of the treatment; It

includes determining the need for medication, evaluating alternative treatment options, arranging the necessary treatment and making appropriate follow-up. The basis of the prescribing process should be the rational decision of the physician, not the pressure made by the patient or the recommendations of the pharmaceutical companies (The Alphabet of Rational Drug Use, 1993, www.e-kutuphane.teb.org.tr).

Rationality of the Prescription Response Process: This process is the point where the pharmacist is involved in the rational use of drugs and his role becomes most evident. Reading the prescription written by the physician correctly and communicating with the physician when necessary, responding to the prescription appropriately (preparing the drugs in a timely manner and presenting them to the patient) is followed by providing appropriate communication, including oral or written education of the patient. In this regard, it is very important to do it without commercial concerns (Yılmaztürk, 2013).

Rational Consumption of Drugs: The parties responsible for rational drug consumption are health personnel, mass media groups, pharmaceutical industries and the most important element, society. In this context, knowing the awareness level of the society and its approaches on this issue play an important role in order for the interventions to be applied to the society to be more effective and appropriate. It is a situation where attention should be paid to raising awareness on health literacy by providing training on disease and drug treatment in the society (Ministry of Health, 2011; The Alphabet of Rational Drug Use, 1993, www.e-kutuphane.teb.org.tr).

Drug Information Support: It is known that drug information must be accurate, complete, impartial, up-to-date, usable and accessible at all stages of rational drug use. It is extremely important to update and inspect the books, brochures, programs, websites, pharmaceutical companies, guides and guides that are sources of information when necessary (Yorulmaz, 2003; The Alphabet of Rational Drug Use, 1993, www.e-kutuphane.teb.org.tr).

Responsibility of the Parties in Rational Use of Medicines

It is known that there are many factors such as lack of education, socio -cultural, economic, administrative and regulatory mechanisms that lead to Irrational Drug Use . There are three main parties in rational drug use: the supplying party, the demanding party and the supervisory-regulatory mechanisms (Ekenler & Koçoğlu, 2016; Sürmelioğlu et al., 2015) (Figure 1).

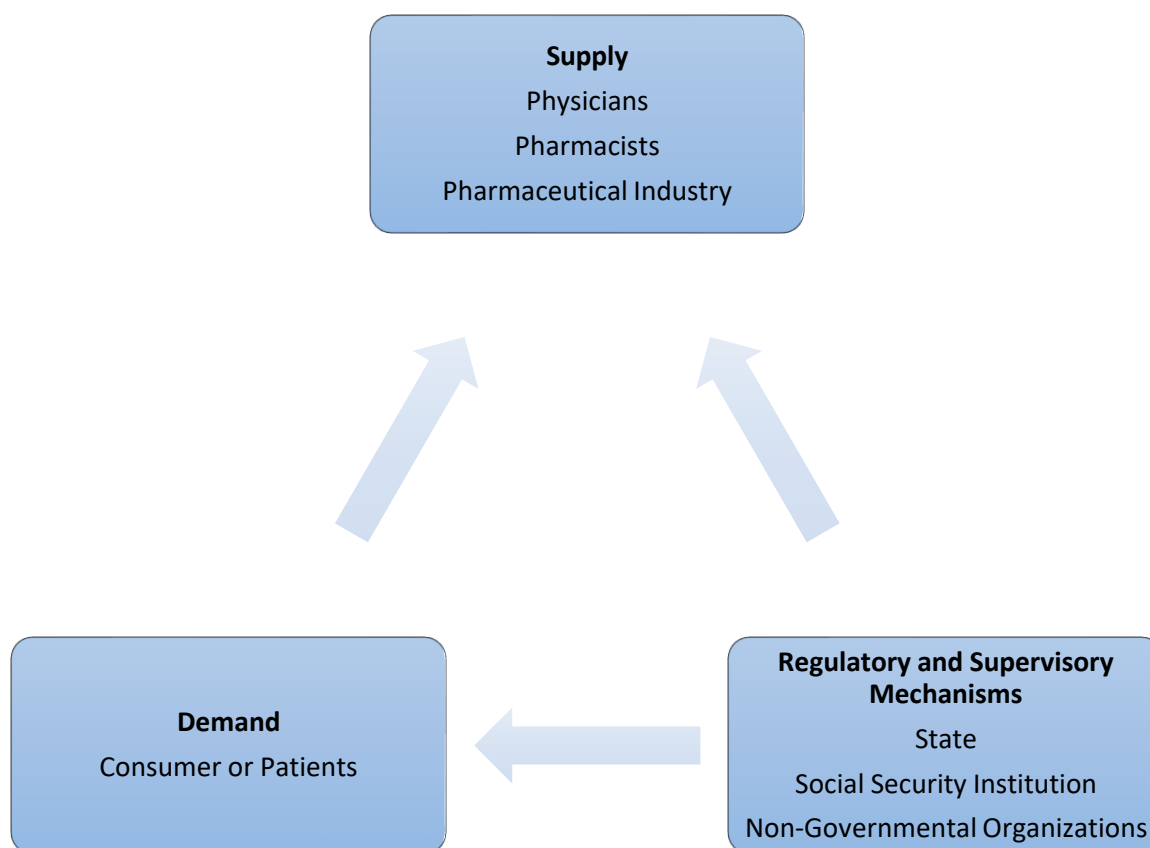


Figure 1. Key stakeholders in rational drug use

Source: (Sürmelioglu et al, 2015)

Health is an important phenomenon for the individual and society. It is also one of the most basic human rights (Nal, 2021). Supply, demand and regulatory mechanisms have responsibilities for rational drug use in health care.

Responsibility of the Physician: Physicians are known as the occupational group with the highest efficiency in rational drug use. The first step of the triangle of physician, patient and pharmacist in rational drug use can be described as a physician (Saygılı & Özer, 2020).

In addition to making a diagnosis, the physician should fulfill the treatment obligation based on the patient's voluntariness in the process of deciding on the treatment method that is compatible with the patient. The physician, who is in close contact with the patient during the communication process, should inform the patient about the results of the diagnosis and treatment of the disease. The physician should show the necessary care and attention during the treatment applied to the patient (Regulation of Medical Deontology 2015).

In general, physicians, who have great responsibilities in the field of drugs and health services in general, need to be trained on rational drug use and their information on the subject should be kept up-to-date (Ministry of Health, 2011).

Responsibility of the Nurse: In the interaction of physician, nurse and patient, the role of the nurse is primarily communication, education and practice. In this context, physicians and nurses act jointly for the same purpose. Drug treatment to be administered to the patient is a process that starts with the decision of the physician and ends in line with the supervision of the nurse. It is extremely important for the society that nurses maintain the necessary care and awareness in the rational drug use process and maximize the benefit to the health system (Ulupınar & Akıcı, 2015).

The nursing regulation, *"In the implementation of the medical diagnosis and treatment plan; to apply the treatments given by the physician in writing, except for emergencies, to accept the verbal medical request of the treating physician in unexpected or sudden situations that develop in the patient and in the diagnosis and treatment plans that must be applied urgently. In this process, patient and employee safety. Statements such as "to monitor the effects of medical diagnosis and treatment interventions on the patient, to keep the necessary records in case of undesirable situations, to notify the physician and to take the necessary measures"* are among the duties of the nurse (TC Resmi Gazete, 2010).

Pharmacist's Responsibility: Pharmacists are responsible for presenting drugs to patients and informing patients about the drugs they use. In the rational drug use process, the second responsibility falls on pharmacists (Barutçu, Tengilimoğlu & Naldöken, 2017).

Although it is the physician who provides the treatment of the diagnosed patient, the last person the patient consults before the treatment is usually pharmacists (Toklu & Dülger, 2011). In addition, pharmacists are the first healthcare professionals consulted when a patient wishes to self-medicate. For this reason, it should guide the patient to the correct use of the drug by supplying the drug in the most correct way (Alpdoğan & Altındış, 2019).

Since the pharmacist is the last step of the treatment process, the role of giving the prescribed drug to the patient, educating the patient about how to use the drug and increasing patient compliance is extremely important in rational drug use (Toklu, 2015).

Patient Responsibility: One of the responsibilities of the society regarding rational drug use is the rational consumption of drugs. This responsibility; should be distributed among the health care workers and mass communicators together with the society (Yorulmaz, 2003).

Rational drug use is the habit that patients acquire during drug use and the process of adherence to the treatment shown accordingly. It is very important to use the prescribed drugs in the amount, time and in accordance with the recommendations made by the physicians. In cases where these items are not fulfilled, the efficiency of other treatment options for the treatments is greatly reduced (Vançelik, Çalkoğlu & Güraksın, 2006). Misuse or abuse of drugs threatens patient safety and leads to waste of resources in this context (Altındış, 2020).

The last element of the quartet, which has the most important obligation in rational drug use, is the patient. The use of the prescribed drugs by the patient in accordance with the recommended dose, duration, interval and warnings is considered as patient compliance. The effectiveness of all steps in rational drug use depends on patient compliance. In order to increase compliance, the physician's allocation of sufficient time to the patient, the education of the patient, the delivery of clearly written drug prospectuses, dose scales, and the reinforcement by nurses in clinics and pharmacists, who are the last factor in pharmacies, have a significant impact (State Planning Organization, 2001).

Regulatory and Supervisory Mechanisms: Many organizations and societies such as the Ministry of Health, health institutions, health personnel, policy makers, reimbursement agency, NGOs and the media are responsible parties for rational drug use (Ministry of Health, 2011). Pharmaceutical industry, which is one of the parties to rational drug use, is also the party that has the obligation to provide health workers with accurate information and appropriate training about their current products and clearly inform them about how to use drugs (Pharmaceutical Industry Employers' Union, 2000).

The stages of reaching the consumer of the drug are production, storage and logistics. All processes are under supervision and control in order to offer safe and effective products to consumers, and all processes are controlled to ensure that the quality standards of drugs produced in the country and imported from abroad are met in a certain amount. All stages in the field of inspection are carried out in Turkey by the *"Turkish Medicines and Medical Devices Agency, Drug*

Inspection Department” (Oktay, 2006).

Family Medicine and Rational Drug Use

In Turkey, where drug use habits are high and the share of drug expenditures in health expenditures is high, the necessity of economical and rational drug use has come to the fore once again with the adoption of family medicine practice (Ministry of Health, 2011).

One of the important indicators in the evaluation of the level of rational drug use of family physicians, who play an important role in the diagnosis and treatment of patients in primary health care services, is the prescriptions offered by the physicians to the patients. Recognition of the contents of the prescription written by the family doctor or being aware of the prescription; It can be shown as a guiding resource in identifying the problems experienced in drug use and eliminating the problems in this regard (Aksoy, Alkan & İşli, 2015). One of the rational drug indicators defined by the World Health Organization is the antibiotic prescription rates of physicians (WHO, 2004).

In this context; Yağız (2020) determined the health literacy levels and knowledge levels of university students studying at Kocaeli University in the 2019-2020 academic year on rational drug use. In addition, the relationship between health literacy and rational drug use levels was examined. As a result of the study, it was determined that 87.6% (n:599) of the students had a high level of rational drug use and 12.4% (n:85) had a low level of rational drug use.

In the researches, it is thought that irrational behaviors are common in the society in rational drug use of individuals, therefore it is thought that more studies on the subject will be positive in terms of public awareness.

Rational Drug Use Practices in Turkey and in the World

Depending on the developments in the world in the current century , studies on rational drug use are carried out in Turkey according to WHO recommendations. In this regard, the “*Guide to good The resource named “Prescribing ”* was translated as “*Guide to Good Prescribing*” by the Ministry of Health in Turkey and the “*Rational Drug Use Program*” was initiated. The guide also covers rational drug use in the light of the basic principles of general drug treatments (Turkish Medicines and Medical Devices Agency, 2011).

Rational Drug Use Branch Directorate was established in 2010 under the Ministry of Health, General Directorate of Pharmaceuticals and Pharmacy. The Turkish Medicines and Medical Devices Agency, the responsible institution for pharmaceuticals, was established on November 2, 2011 (www.resmigazete.gov.tr, KHK/663). Again, in 2011, a study called “*Rational Prescribing Approach of Physicians*” was conducted by the School of Public Health affiliated to the Refik Saydam Hygiene Center Presidency of the Ministry of Health (Turkey Rational Medicines Bulletin, 2015).

“*Regulation on Classification of Medicinal Products for Human Use* ” that prescription drugs can only be obtained under prescription within the scope of the rationality of the prescription fulfillment process . Prescription drugs within the scope of this regulation are clearly stated as: “*Even if they are used correctly, they pose a direct or indirect health hazard if they are not used under medical supervision, and they are frequently and widely misused, and as a result, they pose a direct or indirect danger to human health*” (Resmi Gazete, 2005, www.resmigazete.gov.tr).

Another study was prepared from the National English Formulary, which includes drug-related formulas, to our British National Formulary - BNF literature in 1999 with a team formed by Prof. Dr. S. Oğuz Kayaalp. The sixth and last edition of the “*Turkish Medication Guidelines*”, which was regularly updated in line with the developments, was published in 2011. (Akıcı, www.tfd.org.tr).

In order to make patients' access to medicines from pharmacies safer, they are monitored in a virtual environment with the "Drug Tracking System". The Pharmaceutical Tracking System and DataMatrix Application was implemented on October 1, 2009 in order to prevent counterfeit medicines by keeping track of the medicines and to prevent counterfeit medicines and to generate data for the policies to be prepared regarding medicines (Pharmaceutical Tracking System, www.its.gov.tr).

In addition to these, Hospital Service Quality Standards; The Guide on Rational Drug Use and the Guide on Rational Drug Use Sessions (issued for drug and product license holders) are published through the Ministry of Health (Turkish Medicines and Medical Devices Agency) in order to involve hospitals in this process and to raise awareness of healthcare professionals and those applying to hospitals. , 2011).

It has found an important research area throughout the country with its studies carried out around the world for thirty years on rational drug use. Activities related to rational drug use started in Turkey in the 1990s. The first step that stands out is *"Problem-Based Rational Treatment Training" in Medical Faculties* It was eliminated with the implementation of the (Akıcı et al., 2011).

"Medium-Term Cooperation Program" that WHO and TR Ministry of Health carried out in coordination starting in 1998, the " Pharmacotherapy Training and Clinical Pharmacology Study Meeting on the Placement of Rational Drug Use Principles in Turkey" was held in 1999 . "Turkish Medicines Guide-TIC" was published in 1999 on the use of physicians and pharmacists. Studies were accelerated in 2003 within the framework of the Health Transformation Program (SDP) (Vançelik, Çalıkoglu & Güraksın, 2006).

There are 12 basic recommendations put forward by WHO with the aim of promoting rational drug use. These recommendations are shown in table 1:

Table 1. WHO's recommendations for the promotion of rational drug use

1. To create a supervisory and regulatory institution that will monitor the policies regarding drug use.
2. Utilizing clinical findings and treatment guidelines
3. Based on priority treatments in the creation of a list of essential drugs
4. Establishment of drug committees in hospitals and certain districts
5. Ensuring that student physicians studying at medical faculties receive pharmacotherapy training
6. Providing continuous medical education in-service
7. Developing feedback and monitoring systems
8.Using unbiased and complete information data about drugs
9. Informing and educating societies about drugs
10. Prevention of inappropriate financial incentives
11. Implementation of mandatory and appropriate regulations
12. Taking necessary precautions by securing the availability of drugs and staff

Source: (Turkey Bulletin on Rational Use of Medicines, 2015; WHO, 2002).

Rational drug use is a very important health problem all over the world as it is in Turkey. The “National Action Plan for Rational Use of Medicines” was implemented between 2014 and 2017 in order to train the society and health workers on rational drug use, to increase their level of knowledge and awareness, and to ensure the coordination of all activities. The National Action Plan includes a total of 99 activities, consisting of six main titles and twenty basic strategic objectives. One of the most notable activities is the Prescription Information System (RBS), where prescriptions written by physicians can be monitored and evaluated. In addition to this plan, the "Rational Drug Use National Action Plan 2018-2022 Workshop " of the "Rational Drug Use National Action Plan 2014-2017" was held in Ankara in 2018 (Ministry of Health, 2017; Turkish Medicines and Medical Devices Agency, 2018).

Irrational Drug Use

All behavior, except for the rational use of drugs, can be described as irrational drug behavior. Irrational drug behavior emerges as a result of patients not getting enough benefit from drugs. This situation has an extremely negative effect on the economies of the countries. In Turkey, too, irrational drug use behavior causes a higher rate of waste of drugs (Ekenler & Koçoğlu, 2016).

Irrational drug use can lead to serious drug costs, leading to many health problems, including economic problems and even death. In order to find solutions to these important problems, all countries should take direct measures with their health policies and raise the awareness of the society on rational drug use (Sürmelioglu et al., 2015).

Irrational drug use has become a serious health problem worldwide and in Turkey. Overcoming this problem is through the rational use of personally defined drugs, following the rules set by the WHO, which are universally accepted (Orhaner & Salgın, 2018).

Reasons for Irrational Drug Use

irrational drug use , and among the main problems generally identified in the studies; The antecedents such as prescribing unnecessary amounts of drugs, using unnecessary expensive drugs, not administering drugs correctly, using more antibiotics than necessary or recommending more injections by physicians are among the causes of irrational drug use (WHO, 2002; Sürmelioglu et al., 2015) . In this context , it is seen that many economic and sociological factors also affect the causes of irrational drug use . It is known that the physician is the basis, but pharmacists, other health personnel, society, non-governmental organizations, the pharmaceutical industry and the state also have important responsibilities (Özata, Aslan & Mete, 2008).

irrational drug use , many factors affect another factor and may complicate the problem. This starts from a low level of education and is based on reasons such as social and economic structure, regulatory and administrative mechanisms (Akıcı et al., 2002).

Unnecessary and inappropriate drug use results are seen as an important public health problem in Turkey and around the world. Irrational drug use may cause decreased adherence to treatment, drug interaction, resistance to certain drugs, recurrence or recurrence of the disease, increased incidence of side effects , and increased treatment costs (Why Rational Drug Use, www.akilciilac.gov.tr/).

Consequences of Irrational Drug Use

Reasons such as irrational drug use, life-threatening danger to the patient, unnecessary consumption of resources, side effects of drugs and the possibility of creating dependency emphasize the necessity of rational use of drugs. If drugs are not used rationally, problems such as disruptions and decreases in patients' adherence to treatment, unexpected results, recurrence of the disease and decreased belief in treatment, increase in the recovery period of the disease, frequent side effects, and increased treatment costs lead to problems. This situation greatly increases the

burden on the economies and health systems of countries. As a result, it causes damage to the trust of the societies in the country's health system (Altındış, 2017; Özçelikay, 2001; WHO, 1990).

WHO makes recommendations about the irrational use of drugs in order to prevent the negative effects of health on the economy. In this context, it is extremely important to inform physicians, people in the pharmaceutical industry, pharmacists and the public about the rational use of drugs among the recommendations (Sağır & Parlakpınar, 2014; Altındış, 2017).

Precautions to be Taken Against Irrational Drug Use

In order to prevent or minimize the problems related to irrational drug use, first of all, rational drug use should be ensured. The main points to consider in this regard are;

- The patient's problem must be clearly defined. Undoubtedly, physicians undertake the most important task in this regard (Things to Be Considered, www.akilcilac.gov.tr/).
- In the correct use of drugs, it is extremely important to inform health workers and the community about this issue (Le Grand, Hogerzeil & Haaijer-Ruskamp, 1999).
- to be taken that countries manage their health policies correctly on the rational use of drugs and that physicians and health workers act in a coordinated and conscious manner in this regard (Orhaner & Salgın 2018).
- It is important to increase the amount of lessons related to the problem in addition to the theoretical curriculum during the faculty education period of the physician candidates in terms of reducing the irrational use of drugs (WHO, 2002).
- In order to prevent unnecessary antibiotic use by patients, pharmacies should be directed to physician control before antibiotic sales.
- The society should be kept up-to-date visually and audibly through the media on the rational use of drugs.
- It is important to inform the public that not all drugs on the market will have the same effect on every individual (Ercan & Biçer, 2019).
- Physicians should be prevented from prescribing new and high-priced drugs and making financial profits.
- Increasing the quality of in-service training programs of healthcare professionals and keeping them up to date play an important role in the rational use of drugs.
- In the event that the drug treatment administered by the patient is not completed, waste drugs collection projects should be encouraged in order to prevent child poisoning, drug waste and drug accumulation habits by ensuring the destruction of drugs (Pınar, 2012).

Rational Treatment Process

The rational treatment process begins with the planning of the treatment process after the physician makes the correct diagnosis of the patient in line with the principles of rational drug use. At the stage of diagnosing the patient, the physician sees the patient as a pillar of the treatment process and directs the treatment in the direction it requires. In this context, it starts the treatment process by determining the individual treatment and drug (Akıcı, 2013).

The rational treatment process specified by WHO consists of the following steps (WHO, 1994).

- Determination of the patient's problem by the physician in accordance with the patient's complaint with the correct diagnosis
- therapeutic purpose. At this stage, the goal is “What do you want to achieve with treatment?” is to answer the question. In order to ensure the continuity of treatment compliance between

the physician and the patient, the treatment method should be shared with the patient in a clear language.

- Individualized treatment with the appropriate treatment method and drug selection determined by the physician. While choosing a personalized medicine, principles such as effectiveness, safety, cost and suitability, which are important for rational drug use, should be considered. At this stage, “What are the non-drug treatment options?” search for an answer to the question. It is a priority to provide treatment only by changing the lifestyle. If the change in lifestyle does not have a positive effect in the treatment process, drug therapy should be applied.
- With the drug treatment method, the physician writes a prescription appropriately for the patient and in this context, warns the patient if there is a special situation. Dosage and duration should be determined according to the patient's compliance.
- The last step of the rational treatment process is the physician's evaluation of the treatment applied. At this stage, “Is there a need for the physician to repeat the treatment process with the patient within the scope of the patient's treatment?” The answer to the question is sought (Akıcı, 2013).

Conclusion

The primary goal in rational drug use is to prevent all the physiological, biological and psychological problems that may arise due to the misuse of drugs, and to try to minimize the social and economic damage as much as possible (Ulusoy & Sumak, 2011). While one-third of the world's population struggles to purchase even essential drugs, the ease of access to drugs by some people without a prescription, and the increase in the amount and types of drugs with each passing day show the necessity of the concept of rational drug use (WHO, 2002).

The most basic purpose in rational use is the correct and conscious use of resources. Irrational use causes waste, workload, reductions in service quality and cost increase (Aba, Torun, 2020). The World Health Organization sees it as an extremely important situation to inform physicians, people in the pharmaceutical industry, pharmacists and the public about rational drug use (Sağır & Parlakpınar, 2014; Altındış, 2017).

Bian et al. (2015) and Demirtas et al. (2018), it has been shown that the level of rational drug use knowledge is higher in young individuals, women, individuals in the high income group, and individuals with high education levels. According to Demirtas et al. (2018), on the other hand, the level of rational drug use was found to be higher in individuals under the age of 30, and it was found to be lower in individuals with regular drug use.

In the study conducted by Kan (2022), it was found that rational drug use differs according to the age of the individuals. In the study, it was observed that the rational drug use scale scores of individuals aged 46 and over were higher than those of other age groups. It was concluded that the presence of chronic disease and regular drug use also affect rational drug use. Considering that age increases the chronic disease state and the associated regular drug use, it can be said that the results of the study are significant. It has been stated that this result may be related to the increase in the incidence of chronic diseases as we age. In addition, it can be considered as an important result considering that 90% of individuals over the age of 65, which is accepted as the old age limit, have at least one chronic disease and 60% of them use at least one drug. In general, the mean score of the rational drug use scale was found to be 38.82 ± 3.90 . A score of 35 and above is considered a good result when evaluated as having knowledge of rational drug use. In addition, this result; It was evaluated as a positive result that individuals did not differ between groups according to their age, gender, educational status, income status, occupation, chronic disease status and regular drug use.

It is important when considering the reasons such as irrational drug use, patient's life danger, unnecessary use of resources, side effects of drugs and the possibility of creating addiction accordingly. If drugs are not used rationally, problems such as disruptions and decreases in patients' adherence to treatment, unexpected results, recurrence of the disease and decreased belief in treatment, increase in the recovery period of the disease, frequent side effects, and increased treatment costs lead to problems. This situation greatly increases the burden on the economies and health systems of countries. As a result, it causes damage to the trust of the societies in the country's health system (Altındış, 2017; Özçelikay, 2001; WHO, 1990).

There are some actions that can be taken to prevent or minimize the problems that occur in irrational drug use. One of the most important of these is that physicians define the patient's problems clearly. Another important issue is raising awareness of the society on the correct use of drugs. Making this awareness by all health professionals, especially physicians, pharmacists and nurses, is important in achieving success. In addition, it can be said that the results can be achieved more easily with the applications to be made within the scope of social marketing.

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Remineralization Efficiency of Herbal Products

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Introduction

Dental caries is one of the most common chronic diseases in the world and might cause tooth loss when not treated properly. The concept of modern dentistry aims to prevent the progression of caries with early diagnosis, and to improve the aesthetics and function of the tooth with the remineralization process. Thus, caries lesions without cavity are treated with a non-invasive method (Nagarathana et al., 2015). Demineralization occurs through the leaving of mineral ions in dental hard tissues from hydroxyapatite crystals. Restoring these mineral ions to hydroxyapatite crystals is called remineralization. Both processes take place on the tooth surface and a large number of mineral ions can be lost from hydroxyapatite without deterioration of tooth integrity. As a result of the deterioration of the hydroxyapatite structure, cavities are formed. Demineralization is a reversible process; therefore, partially demineralized hydroxyapatite crystals in teeth can be remineralized (Higham, 2016). Enamel caries diagnosed at the initial stage can be treated by regulating diet, controlling plaque formation, and applying appropriate antibacterial agents, thus, demineralized dental tissue can become remineralized (Guerrieri et al., 2012).

Formation of Initial Enamel Caries

The caries formation process is a gradual process that requires periods of repeated and prolonged exposure of the enamel to acidic attacks, with the environment dropping below the critical pH (pH 5.5) and returning to the intermediate medium's plaque pH (pH 7.0). In the development of dental caries, the stage before cavitation is called the initial caries lesion. It is characterized by subsurface demineralization areas that occur under an intact enamel surface (Kidd & Fejerskov 2004) (Figure 1). The mineral content of the affected area is reduced, which affects the translucent nature of enamel and the color of these areas appears more opaque white. For this reason, initial enamel lesions or flat surface caries are also called white spot lesions and are the first visible signs of caries formation (Øgaard et al., 1988). It is important to distinguish the etiology of these lesions clinically, because white lesions can be caused by caries or be hereditary. When the tooth surface is air-dried, caries lesions appear opaque and chalky. This appearance is due to the loss of its translucent feature due to subsurface demineralization of the enamel tissue in this area. In contrast, developmental lesions are less or unaffected by air drying (Fejerskov et al., 2015).

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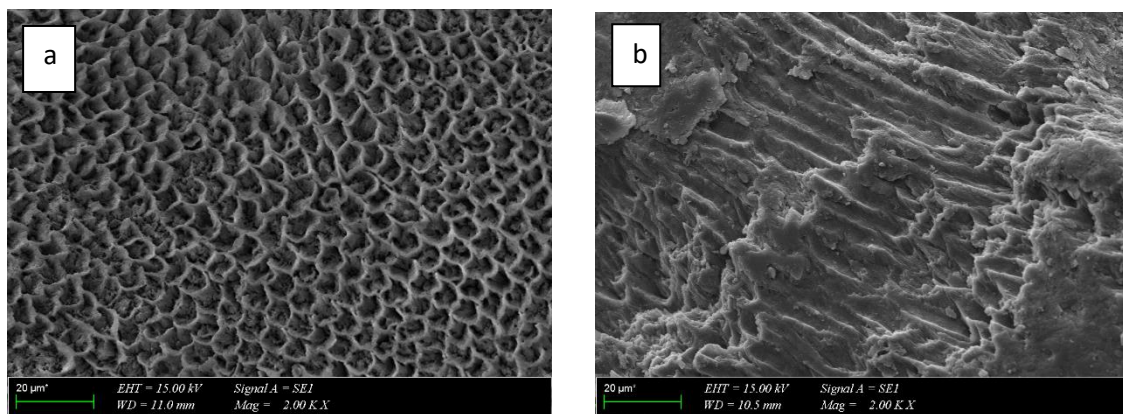


Figure 1. *Demineralized enamel tissue: a) coronal section b) sagittal section.* (Alkan, 2022)

Treatment of Initial Carious Lesions

Various options are available for the treatment of initial carious lesions. Recent approaches are listed in the following sections.

Providing Oral Hygiene

Dental caries is an important problem that affects not only the individual, but also social institutions and the health economy. Untreated caries affects millions of people worldwide. Such a big problem also causes huge cost. Raising awareness about individual oral hygiene and daily brushing with fluoride toothpaste is one of the solutions for tooth decay. Providing personal oral hygiene with tooth brushing and flossing, removing the bacterial plaque and thus changing the initial carious lesion formation process are the most effective methods that patients can apply individually (Hicks et al., 2005).

Diet Regulation

Apart from the presence of bacterial plaque, another factor affecting caries formation is diet. The foods taken can act as an inhibitor of caries formation and can be among the causes of caries formation. For example, some foods with a hard and fibrous structure help mechanical cleaning, while others show a caries preventive effect by increasing the amount and flow rate of saliva with their taste and smell. Food and beverages such as cheese, milk, cocoa, tea and proteins have a bacteriostatic effect by changing the metabolism of cariogenic bacteria. However, fermented carbohydrates can cause caries when consumed frequently (Kalender, 2017).

Use of Antimicrobial Agents

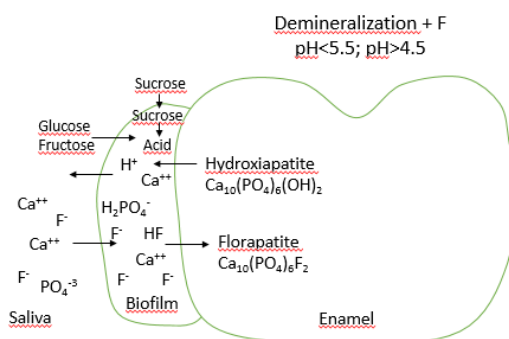
The use of chlorhexidine was first offered to the public in the USA as a 0.12% mouthwash for periodontal treatment of high-risk patients. It reduces the number of *Streptococcus mutans* and accelerates remineralization. It is recommended to use a chlorhexidine mouthwash for 30 seconds just before going to bed, as the reduced salivary flow rate at night helps chlorhexidine to bind to the structures in the mouth more easily. It has been reported that when chlorhexidine mouthwash is used in this way for 2 weeks, it reduces the number of *Streptococcus mutans* below the potential for caries formation, and the effect of this reduction lasts between 12 and 26 weeks (Roberson et al., 2006).

Remineralization of Initial Carious Lesions

Initial caries lesions can be remineralized by preventing the factors that cause caries formation and increasing the duration of the protective factors in the mouth. In general, remineralization is a natural repair process and many methods have been proposed to improve it. The dynamics of formation and prevention of the initial caries lesion occurs between demineralization, which is the process of removing mineral ions from the hydroxyapatite crystals of enamel, and remineralization, which is the process of restoring these lost mineral ions, particularly calcium and phosphate ions, into the cavities in the hydroxyapatite crystals (Figures 2,3).

An ideal remineralizing agent (Walsh, 2009):

- It should provide optimum level of calcium and phosphate ions, thus increasing remineralization and preventing the formation of tartar.
- It should be able to penetrate deeply into the subsurface demineralized area.
- It should be in active form even below the critical pH and thus prevent demineralization.
- It should have antimicrobial activity against cariogenic microorganisms.
- It should show a synergistic effect with the remineralization capacity of saliva.
- It should be effective even in cases where there is a decrease in salivary flow rate and in xerostomia patients.
- It should not cause any negative effects.
- It should be easily accepted by the society.



Tenuta 2008).

Figure 2. Enamel demineralization in the presence of F in dental biofilm. Carbohydrates (sucrose, glucose, fructose) are converted into organic acids by bacteria in the biofilm. When the pH drops below 5.5, the saturation of hydroxyapatite (HA) in the biofilm decreases, causing mineral dissolution. However, if the pH is higher than 4.5 and F is present, the biofilm Florapatite (FA) becomes supersaturated and mineral re-precipitation occurs in the enamel. As a result, demineralization is reduced (Cury &

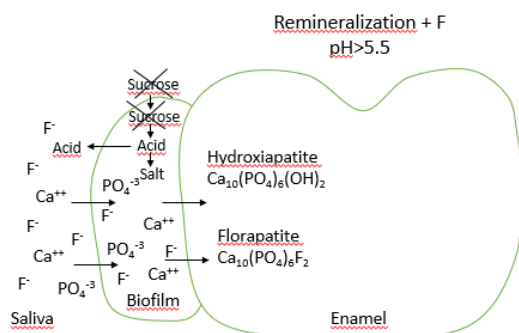


Figure 3. Enamel remineralization in the presence of F in dental biofilm. After the intake of carbohydrates has ceased, the acids in the biofilm are cleared by saliva and converted into salts. As a result, the pH increases and at 5.5 or higher the biofilm becomes supersaturated with HA and FA. Thus, Ca and P lost by the enamel can be recovered more efficiently if F is still present in the biofilm (Cury & Tenuta 2008).

Fluoride Applications

Fluoride shows its effect against caries by three different mechanisms. First, the presence of fluoride greatly enhances the formation and accumulation of fluorapatite, which is formed by the combination of calcium and phosphate ions in saliva. Fluorapatite is resistant to dissolution; therefore, it replaces salts containing manganese and carbonate, which are easily dissolved and lost due to demineralization, making the enamel more resistant to acids. The second mechanism is remineralization of the initial cavity lesion with fluorapatite crystals. The third and final mechanism is the antimicrobial activity of fluoride ions. Low fluoride concentrations inhibit the production of the glycosyltransferase enzyme. The glycosyltransferase enzyme enhances bacterial adhesion and provides glucose for extracellular polysaccharide formation. Topical fluoride applications at high concentrations (12,000 ppm) have a direct toxic effect on oral microorganisms, including mutans streptococci (Roberson et al., 2006). Fluoride applications can be classified as systemic and topical.

Systemic Applications

Systemic applications are effective methods, especially in individuals with a high risk of caries formation and in societies where fluoride use is low. Fluoridation of water, which is one of the systemic applications, shows not only systemic but also topical effects (Roberson et al., 2006). The World Health Organization (WHO) has reported that consuming 1 mg of fluoride per day is beneficial for health. Systemic applications of fluoride are the fluoridation of drinking water, salts and milk, and the addition of fluorinated tablets or drops to the diet.

Topical Applications

Systemic fluoride intake was thought to be effective for a long time before the application of fluoride directly to the tooth surface was initiated. However, it is now accepted that the use of topical fluoride is more beneficial during tooth development and maturation. Topical fluoride in high concentrations (12,000 ppm), often professionally applied in dental clinics, has a direct toxic effect on oral microorganisms, including *Streptococcus mutans* (Roberson et al., 2006). Fluoride varnishes have been developed to prolong the contact time with the tooth, to hold onto the enamel for a longer time, and thus to prevent rapid fluoride loss after application. Varnishes act as a reservoir for slow release and facilitate further fluoride uptake (Øgaard et al., 1994). In the guidelines of the American Academy of Pediatric Dentistry, it is recommended to apply fluoride varnishes at a concentration of 5% (22,600 ppm) at least twice a year for primary teeth and two or four times a year for permanent teeth (de Sousa et al., 2019).

Casein Phosphopeptide–Amorphous Calcium Phosphate (CPP–ACP) Applications

ACP is a tricalcium phosphate containing calcium and phosphate ions in an amorphous structure. When ACP enters a solution, it rapidly transforms into a stable structure such as octacalcium phosphate or apatite. Casein is a phosphoprotein that makes up 80% of the proteins found in cow's milk. The most important feature of casein is the retention of calcium and phosphate ions in protein complexes. These ions break down into smaller peptides like CPP, making them highly durable. Casein molecules act as a carrier, providing calcium and phosphate ions that tissues such as teeth or bones can use for remineralization. Mendes et al. reported that the use of CPP-ACP is a good alternative for remineralization of initial carious lesions (Mendes et al., 2018). The remineralization effect can be improved when this product is applied together with fluoride.

Nanohydroxyapatite

Today, nanotechnology is developing by offering various caries preventive and therapeutic applications. Nanotechnology has sought to replicate the nanostructural properties of human

enamel and offer biology-inspired strategies for remineralization and caries treatment (Chandki et al., 2012).

In dentistry, nanoparticle application is used in two ways as preventive dentistry and restorative dentistry (Maman et al., 2018). In recent years, various hydroxyapatite types of calcium carbonate in nanometric sizes have been applied in the prevention and treatment of incipient caries lesions. In dentistry, the sizes of hydroxyapatite nanocrystals vary between 50-1000 nm; being so small in size increases the surface area they can bind to, so they can bind to proteins, biofilm and bacteria when present in the toothpaste. Likewise, thanks to their small size, they can repair small cavities and indentations in the enamel surface. Nanohydroxyapatite, a bioactive and biocompatible material, contributes to the remineralization process with high amounts of calcium and phosphate ions from oral fluids, increasing crystallization by directly filling the micropores of the initial caries lesions. It has been reported that a 10% nanohydroxyapatite concentration is optimal for remineralization of initial enamel caries (Huang et al., 2009).

Sodium Calciumphosphosilicate (Bioactive Glass)

When bioactive glass comes into contact with saliva, it rapidly releases sodium, calcium and phosphorus ions available for remineralization of the tooth surface into the saliva and increases the pH. The released ions directly form hydroxycarbonate apatite (HCA). They also attach to the tooth surface and continue to release ions and remineralize the tooth surface after the first application. These particles have been shown to convert to HCA for up to 2 weeks by releasing ions (Du et al., 2004). Novamin, a commercial form of bioactive glass, adheres to the exposed dentin surface and forms a mechanically strong and acid-resistant mineralized layer. There is a sustained release of calcium that maintains its protective effects on dentin over time (Burwell et al., 2010).

Natural Remineralization Systems

Plant extracts have been used as medicine in traditional medicine since ancient times. Today, these plant extracts are of particular interest because they are not chemical or synthetic. If the effect of herbal products is supported by evidence as a result of scientific-based studies, they may be better and safer alternatives for treatment (Shaheen et al., 2015). In recent years, the desire to return to natural products has also affected the view against the use of herbal agents as antimicrobial agents in dentistry. Antimicrobial properties of essential oils obtained from plants instead of artificial molecules are being investigated. Besra et al. investigated the antimicrobial properties of various plant extracts on caries pathogens and concluded that the plants can be used alone or in combination in the treatment of dental caries (Besra & Kumar 2018). Studies have been conducted in which herbal extracts are used as antimicrobial plaque agents to prevent dental caries and reduce gingivitis (Poureslami, 2012).

Some natural plant extracts have antimicrobial properties and have the potential to prevent demineralization and provide remineralization of enamel (Philip, 2019). Among the remineralization systems, fluoride-free and natural systems will reduce the risk of fluoride toxicity. Because plant-based phytochemicals and bioflavonoids can provide remineralization of the initial caries lesion, they will be more acceptable to the public than artificial chemical derivatives and fluoride-based systems used for dental caries prevention. Common oral health problems such as dental caries, periodontitis, microbial dental plaque and gingivitis can be prevented and treated if appropriate herbal treatments are used consistently over a period of time (Bilgin Göçmen et al., 2020). Also, most of them are natural foodstuffs that show no toxicity and are "generally recognized as safe" (GRAS) by the US Food and Drug Administration (FDA) (Annamalai et al., 2020). Some of the natural and herbal extracts are listed below.

Cocoa Bean

Theobromine is an alkaloid found in cocoa beans (*Theobroma coca*) and chocolate. It is a white crystalline powder from the methylxanthine family (3,7 dimethylxanthine). Theobromine has been shown to increase crystal growth, resistance to acid attack, and microhardness of enamel (Amaechi et al., 2013; Syafira et al., 2013). In another in vitro study, it was reported that theobromine has a remineralization capacity similar to that of fluoridated toothpastes (Taneja et al., 2019).

Xylitol

Xylitol is a naturally occurring five-carbon sugar polyol (pentylol) found naturally in fruits, vegetables, and berries. It is mainly derived from xylan-rich plant materials such as birch and beech (Bar, 1988). It is used instead of sugar. It has been reported that the use of chewing gum containing xylitol increases the salivary flow rate, improves the protective properties of saliva and helps remineralization. It has been suggested that caries-associated mutans streptococci are unable to use xylitol in their metabolism and therefore reduce acid formation in the oral biofilm. It has been reported that xylitol significantly reduces the incidence of caries and increases tooth remineralization (Hildebrandt & Sparks, 2000; Ritter et al., 2013).

Ginger

Ginger (*Zingiber officinale*) is one of the most widely used herbs in traditional medicine. Numerous pharmacological activities such as antioxidant, antibacterial, anti-inflammatory, antinociceptive, antimutagenic and hepatoprotective have been shown in many studies (Abdel-Azeem et al., 2013; Jeena et al., 2013; Jeena et al., 2014). In addition, there are studies showing the antibacterial effect of ginger against many microorganisms, including mutans streptococci (Fatima et al., 2000; Khan et al., 2010; Islam et al., 2012).

Ginger content; It contains approximately 50% carbohydrates, 9% protein and free amino acids, 6-8% fatty acids and triglycerides, 3-6% ash and 3-6% crude fiber (on the basis of dry matter), although it may vary depending on the type and geography of cultivation (Tang & Eisenbrand, 1992). Ginger is a good source of essential micronutrients such as potassium, magnesium, copper and manganese. Ginger rhizome also contains small amounts of vitamins A, E, and some B- and vitamin C (PR & Prakash, 2010). Ginger is a non-toxic natural product and is considered "generally safe" (GRAS) by the U.S. Food and Drug Administration (FDA). The pungent oil component in it consists of a series of polyphenolic ketones with many pharmacological activities (Ravindran & Babu, 2016). The bioactive components gingerol and shagelol obtained from the ethnolic extracts of ginger exhibit antifungal and antibacterial activity. It is thought that ginger's remineralization capacity is due to its antimicrobial properties and high fluoride content (Al-Duboni et al., 2013).

Ginger is one of these herbal extracts with its antibacterial properties. Ginger essential oil and oleoresin have shown significant antioxidant and antimicrobial activities (Bellik, 2014). Ginger; It has been included in oral and dental care products as an accepted and renewable natural product due to fewer side effects, better patient tolerance, relatively cheaper, long history of use (Ody, 2000; Al-Achi, 2001; Abebe, 2003; Barrett, 2004).

Bilgin et al. conducted two studies with a herbal mixture consisting of chocolate, ginger and honey, and in both in vitro and in situ studies, they showed that tooth cream containing ginger and honey had very positive effects on remineralization as a result of both microhardness and fluorescence evaluations compared to alternative normal fluoride toothpaste. (Bilgin et al., 2016; Bilgin Göçmen et al., 2016).

Rosemary

Rosemary (*Salvia rosmarinus*) is a small aromatic shrub composed of terpenoids, flavonoids, phenols and essential oils with antimicrobial, antifungal and antioxidant properties. Al-Duboni et al. reported that methanolic extract of rosemary (30g/100mL) had inhibitory effects on mutans streptococci, and stated that rosemary extract was effective in remineralization as a result of fluorescence and microhardness evaluations on enamel (Al-Duboni et al., 2013). Bilgin et al. reported the in vitro remineralization capacity of a mixture of rosemary, ginger extract and honey (Bilgin et al., 2016).

Hesperidin

Hesperidin is a member of the flavanone group of flavonoids isolated in large quantities from the peels of citrus fruits such as lemons, oranges, grapes and tangerines. Hesperidin has been shown to have antioxidant, anti-inflammatory and anti-carcinogenic properties. It has been reported that Hesperidin inhibits demineralization, increases remineralization in dentin, and preserves collagen even under fluoride-free conditions (Islam et al., 2012; Li & Schluesener 2017).

Grape Seed Extract

Grape (*Vitis vinefera*) seed extract contains a powerful antioxidant polyphenol, proanthocyanidin. It strengthens collagen-based tissues by increasing collagen cross-links. Proanthocyanidin has been shown to inhibit root caries by interacting with microbial cell membrane proteins and lipids, leading to cell membrane disruption. Proanthocyanidin inhibits the enzymatic activity of ATPase and amylase. At the same time, it inhibits the glucosyl transferase enzyme produced by mutans streptococci, preventing bacterial adhesion to the tooth surface, which shows anticariogenic properties. It has been shown that grape seed extract can be used as a substitute for fluorides in the prevention of root caries in elderly patients. An in vitro study reported that mouthwash containing grape seed extract limits dentin matrix degradation and increases remineralization. A recent systematic review concluded that the caries prevention of grape seed extract compared with fluoride inhibits the proliferation of bacterial biofilms on the tooth surface and promotes remineralization (Xie et al., 2008; Khaddam et al., 2014; Delimont & Carlson, 2020).

Galla Chinensis

Galla chinensis is a traditional Chinese herb that has been thought to have anti-caries effect in recent years. It has been stated that *Galla Chinensis* prevents demineralization and contributes to remineralization and increases the effect of fluorine. According to a hypothesis that emerged as a result of the studies, they stated that *Galla chinensis* can affect the remineralization of the caries lesion, and that it does this by combining with the organic matrix in the enamel. The possible effect of *Galla chinensis* is thought to be through its polyphenols, which combine with the organic matrix in the enamel and block the ion diffusion pathways. Polyphenol compounds have been reported to act as calcium ion carriers to the lesion body (Cheng et al., 2010; Zhang et al., 2016; Philip, 2019).

Honey

Honey is a natural food product produced by honey bees (*Apis mellifera*), consisting of nectar secreted from flowers and substances secreted by insects. Due to its chemical composition, honey has been used for many years in the treatment of burns and infected wounds (Medhi et al., 2008; Mandal & Mandal, 2011). Honey has been found to have promising potential as an alternative agent in the management of disease conditions such as dental caries and gingivitis following orthodontic treatment (Atwa et al., 2014).

Honey, one of the agents traditionally used with herbal medicines, has a bacteriostatic effect on pathogens and has a pH of about 3.9. Studies have shown its effectiveness on oral pathogenic

bacteria (Al-Hasani, 2018; Sateriale et al., 2020). Honey has antibacterial effects on *S.mutans*, *L. acidophilus*, *A.viscosus*, *P.aeruginosa*, *V.alcaligenes* and *S.aureus*. Honey has a pH of 3.9 so it is acidic and can inhibit the growth of pathogens as most pathogenic microorganisms thrive at pH 4.0-4.5. Diluting honey with saliva will increase the pH and reduce its effect. This contributes to its antibacterial activity by increasing the enzyme activity and the effect of glucose oxidase enzyme and the production of hydrogen peroxide, which is an oxidizing agent (Molan, 1992; Brudzynski et al., 2011). The glucuronic acid produced by glucosoxidase in honey helps calcium absorption (Eteraf-Oskouei & Najafi, 2013). In addition, it has been reported that orally taken honey reduces prostaglandin levels and has prebiotic and antibacterial effects (Al-Waili et al., 2011; Al-Waili et al., 2011).

Conclusion

Although modern chemicals are widely used today, the demand for herbal/natural products has increased in recent years. Plant extracts, which are frequently used in traditional medicine, gain importance as an alternative treatment for dental health. As a result of the researches and achievements in the field of herbal medicines and formulations, many problems of oral health might be prevented and treated. Today, there are many herbal medicines and derivatives, toothpastes, mouthwashes, etc., which are related to oral and dental health, sold with or without a prescription exist. Natural products are potential and valuable resources for the development of medicines and products to manage oral diseases. Although some progress has been made in this area, there is still limited information about the safety and effectiveness of these products. Thus dentists should be skeptical about the utilization of products that do not have sufficient scientific studies and proven efficacy, and should pay attention not to mislead the society.

Further clinical research is needed to develop drugs and products containing active plant extracts or phytochemicals that can be offered as a supplement to existing treatment or as an alternative treatment. Dentists should be knowledgeable about these products as they are natural, safe and economical. They should also choose to prescribe and recommend these products in their treatments.

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The Relationship of Exapolysaccharides with Biofilm and Quorum Sensing

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Introduction

Probiotics, derived from the Greek and meaning “for life”, is a term used to define living non-pathogenic organisms which have a demonstrated beneficial effect on the host when ingested in adequate amounts (FAO/WHO, 2001). The most commonly used probiotics are represented by lactobacilli, bifidobacteria and non-pathogenic yeasts. They must be capable of surviving the passage through the stomach to enter the intestinal tract, where they exert helpful effects on the gut microbiota, and of resisting to the stressful environments during industrial processes. The most important and known beneficial effects of probiotics include the prevention of diarrhea, constipation, and food allergies; other factors, such as a reduction of gas production, changes in bile salt conjugation, anti-bacterial, anti-inflammatory and anti-viral effects, may result relevant to the benefits of specific probiotics (Barrett et al., 2008).

Biofilms and Quorum Sensing

Biofilms form a heterogeneous matrix consisting of microbial communities and their secreted extracellular polymeric substances (EPSs) (Wingender et al., 1999) and in sediments incorporate particles within the matrix. Microorganisms that inhabit sediments and form biofilms are highly adapted for this habitat and secrete adhesive polymers. The EPS is primarily composed of organic molecules such as polysaccharides, proteins, lipids, and nucleic acids (Stoodley et al., 2002). EPS affects sediment dynamics by, directly enhancing the cohesive forces between sediments, binding them together, and coating sediment particles and changing the micromorphology of individual grains (Paterson et al., 2009). Bacteria in natural habitats predominantly exist as matrix-encased species-rich communities, either as surface associated biofilms or surface independent floccular aggregates. This is in contrast to their planktonic single species population counterparts commonly studied in laboratory conditions. The formation and dispersal of structured bacterial biofilms or aggregates occur in response to a range of environmental cues and signals, such as changes in nutrient concentrations, oxygen, temperature, as well as chemicals and predatory stresses (Matz et al., 2004).

Bacterial biofilm formation is often regulated by quorum sensing (QS) which is a population density-based cell-cell communication process via signaling molecules. Numerous signaling molecules such as acyl homoserine lactones, peptides, autoinducer-2, diffusion signaling factors, and α -hydroxyketones have been studied in bacteria. Genetic alteration of QS machinery can be useful to modulate vital characters valuable for environmental applications such as biofilm formation, biosurfactant production, exopolysaccharide synthesis, horizontal gene transfer, catabolic gene expression, motility, and chemotaxis (Fig 1). These qualities are imperative for bacteria during degradation or detoxification of any pollutant. QS signals can be used for the fabrication of engineered biofilms with enhanced degradation kinetics (Mangwani et al., 2016).

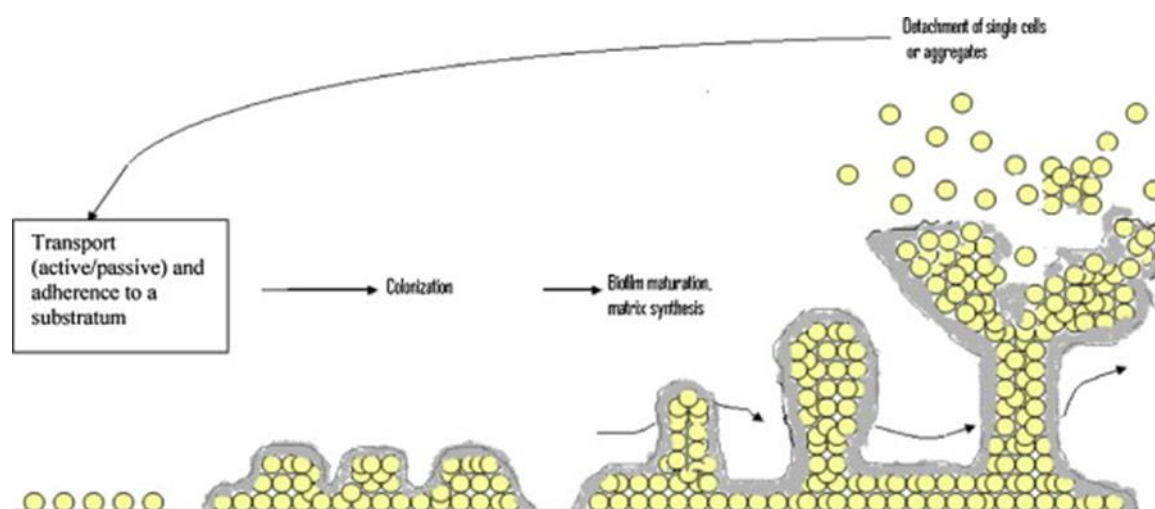


Fig. 1. The cyclic evolution of a biofilm at cellular level, from single cells, sedimented and adherent to a conditioning pellicle on a substratum, cellular multiplication and colonization of substratum, formation of a multilayer biofilm with columns and mushroom-like structures, and finally the detachment of single cells or aggregates and their dissemination (Lazar, 2011).

QS also regulates many biological behaviors of the mature biofilm. For example, the expression of several extracellular virulence factors, including exotoxin A, elastase, pyoverdine and rhamnolipid (Kirisits and Parsek, 2006). QS factors are critical to establishing infection in the host and contribute antibiotic resistance properties, as well; therefore, inhibition of the QS system is believed to be a promising strategy to attenuate production of virulence factors and protect against infection. Secretion of an extracellular polysaccharide substance that acts as a physical protection barrier for the encased bacteria. There is a pressing demand for targeted therapies that are capable of killing cancer cells selectively without affecting normal healthy cells or at least to act as adjuvants to lower the therapeutic doses and increase efficiency of conventional anticancer drugs. Synthesis of EPSs by LAB is a well-known phenomenon on which exists as a cell-bound EPS, adhering closely to the bacterial surface, or releases EPS into the surrounding medium (Paulo et al., 2012). The polysaccharide structure may affect bacterial adhesion to mucus. Other authors have claimed that, in some cases, the EPS envelope covering the producing strains hinders bacterial adhesion (Nikolic et al., 2012).

Lactobacillus EPSs have been reported to contain several functional groups, for instance, carbonyl, phosphate, and hydroxyl groups, which were suggested to play an essential role in exerting the antimicrobial and antioxidant effects of EPS (Riaz Rajoka et al., 2020). These polymers showed in vitro or in vivo inhibition of Gram-positive and Gram-negative pathogens with different degrees of resistance. The Gram-positive bacterial cell wall includes numerous structural components, which are essential components involved in the interaction between bacterial cells and various receptors on other surfaces (AlKassaa et al., 2014). The cell surface of bacteria is a vital element in cell-to-cell and cell-to-host communications (Dertli et al., 2015). Many of the studies reviewed in this report did not examine the electrical charge on the polymer. For instance, negatively charged EPS from the *Lactococcus lactis* F-mou strain revealed higher inhibitory action against Gram-positive than Gram-negative pathogens, with *Bacillus cereus* ATCC 10702 demonstrating the greatest inhibition (Nehal et al., 2019). There is no definitive mechanism to explain the antibacterial

action of the EPS produced by the LAB against Gram-positive and -negative bacteria. Attempts to investigate a potential mechanism(s) to explain the observed antibacterial activity are continuing. It was thought that because EPS was able to disrupt the structure of the bacterial cell envelope, especially the peptidoglycan layer, it was proposed as a potential inhibitory mechanism (Sivasankar et al., 2018). These biofilm communities containing beneficial organisms such as lactobacilli may play an essential role as protective agents (Aoudia et al., 2016).

Biofilm formation is a complex and dynamic process comprising five steps including: initial attachment, irreversible attachment, initial growth of the biofilm structure, maturation, and dispersion (Spanò et al., 2016). Bacteria release chemical autoinducers (AIs) through the action of QS that accumulate within the bacterial environment (Fig 2). Through detecting changes in AI concentrations, bacteria exchange information. A range of signaling molecules related to bacterial QS has been discovered. Dependent on the type of signaling and sensor systems, the QS of bacteria can be classified, we summarize the pathways associated with each QS system and its distribution in bacteria (Ran et al., 2016). Inhibition of signaling molecule generation: the QS process is inseparable from the generation and participation of signaling molecules. By inhibiting related enzymes in the signaling molecule synthesis pathway, the generation of signal molecules can be blocked and QS can be inhibited. For example, Triclosan can inhibit enoyl-ACP reductase an important protein in the acyl-homoserine lactone (AHL) generation process (Fischer et al., 2015).

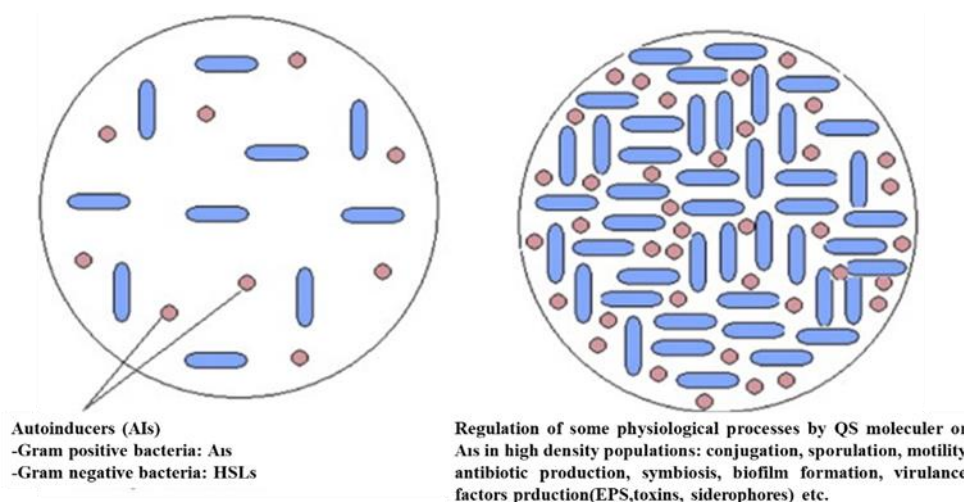


Fig 2. Cell to cell communication by QS mechanism, mediated by small molecules or autoinducers (Lazar, 2011).

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The Problems Experienced By Infertile Couples and the Nursing Approach

Fatma BAŞAR¹

Introduction

Infertility is defined as the inability to conceive or maintain a pregnancy despite having regular sexual intercourse (two or three times a week) for at least one year without using any contraception method. It is reported that 10-15% of couples of reproductive age in the world have infertility problems and more than 80 million people are affected by infertility. In our country, it has been determined that the rate of married women between the ages of 15-49 who have no children and who are not able to have children is 11.2%. According to the 2018 data of the Turkey Demographic and Health Survey (TNSA), the rate of married women between the ages of 15-49 who have no children and stated that it is not possible to have a child is 5%. Approximately 40% of infertility cases, which is one of the important questions of our age and affects many couples, are women, 40% are men, and the remaining 20% are combined causes.

Infertility diagnosis and treatment process is a difficult and long process that affects couples physically, psychologically and financially. In this process, couples may experience stress, panic and intense emotional pressure. At the same time, in societies where femininity is equivalent to motherhood and men's productivity, fertility is accepted as an indicator of an individual's sexual identity. The society's ostracism of the woman without children and the questioning of her husband's masculinity may negatively affect the mental health and quality of life of the couple, leading to depression, anxiety, decreased sexual desire and deterioration in marital relations. In many studies, it has been determined that infertile couples experience psychological, social and sexual problems. Although both spouses are affected during the diagnosis and treatment of infertility, women feel more stress, guilt and responsibility than men due to the effect of interventions on the female body.

In this section, it is aimed to understand the problems experienced by infertile couples by examining the socio-cultural, psychosocial, economic, sexual problems and quality of life of infertile couples in line with the literature.

Sociocultural Problems in Infertile Couples

Marriage brings with it having children and problems may arise in marriages where there is no child. Birth not only increases the respect for the woman, but also strengthens the place of the woman in the family, relatives and group. The father, on the other hand, not only looks to the future with confidence, but also gains prestige among his friends and relatives. Because no matter how despised the infertile woman is by her relatives because she cannot give birth, the man also feels the social and mental oppression of the pressure coming from the environment and the 'not being put in the place of a man'. A woman who does not have children is not treated well in the family, and a woman who is called "infertile" can be described as "unlucky". As in many countries, the words woman and mother are used synonymously in Turkey, and childlessness in such a

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cultural structure is quite a degrading situation for women. In our country, couples who do not have children in rural areas are excluded by the society, but more blame falls on the shoulders of women, and the woman has to accept the marriage in order for her husband to have a child. One of the basic instincts in humans is the reproductive instinct. It may become inevitable for sexuality to go beyond the reproductive function, to experience feelings of failure and inadequacy at this stage, and to create problems that spread to many areas of life. Inability to reproduce often creates a social stigma and is perceived as a shameful inadequacy.

Cultural, ethnic or religious factors also impose new burdens on the infertile couple. For example, there is a lot of superstition and misinformation about infertility, especially in vitro fertilization (IVF) treatment in Turkish culture. Kamaç (2003) found in his study that 28% of women and 8% of men visit the *hodja-mausoleum-çeşme*, 7% of women and 36% of men use herbal medicine, and about half of the couples resort to traditional methods.

Childbirth is perceived as a divine reward, while infertility is perceived as divine punishment. In some religions, assisted reproductive techniques are considered a sin, and if the woman is infertile, her marriage is perceived as inadequacy.

Psychosocial Problems in Infertile Couples

Although the reactions of individuals to infertility show individual differences, it is stated that there are similar aspects. For couples, the common feelings of not being able to have children are the feeling of frustration and the lack of parental roles in society.

Since infertility is a multidimensional problem, it negatively affects the psychological well-being of couples. D'Souza et al., in their study, stated that the psychological well-being of women who received infertility treatment was low. In Klemetti's study; states that the psychological well-being of infertile women and men is very low and that this situation is related to anxiety and depression. In addition, it has been reported that the emotional stress experienced during the diagnosis and treatment of infertility may cause couples to think that they cannot have children and lead them to a sense of hopelessness. Kaya et al., in their study, found that infertile women experience stigma and hopelessness. determined that hopelessness increased as social support decreased in infertile women.

Depression and anxiety levels were compared among infertile couples in Poland in 2009. According to Beck depression scale; Depression level was found to be 28.6% in infertile women, 18.4% in fertile women, 9.0% in infertile men, and 11.0% in fertile men (Drozdol and Skrzypulec 2009). In the study of El Kissi et al. (2013) with couples who received infertility treatment, it was determined that women experience more anxiety and depression than their spouses. In addition, women's self-esteem was found to be lower than their spouses [16]. In Yusuf's study (2016), it was found that 70% of infertile women experience varying degrees of anxiety and 58% experience moderate-to-severe anxiety.

Psychosexual Problems Experienced by Infertile Couples

Evaluation of infertility is a condition that is emotionally disturbing and also sexually disturbing to individuals. For example, when taking the history of the couples, the couples are asked questions about their sexual performance such as copulation techniques, sexual desire and responses, history of abortion, discomforts during intercourse, knowledge of sex and reproduction. Such questions may be seen as threatening, disturbing and inappropriate by couples. In some cases, questions focusing on sexual competence can lead to sexual failures or changes in sexual behavior over time.

Infertile couples frequently experience sexual problems such as sexual reluctance, anorgasmia, decreased sexual satisfaction, and dyspareunia as sexuality becomes an act to have children. In the study of Tao et al. (2011) it was stated that infertility affects the sexual activities of

individuals, and there is a tendency to decrease in the frequency of sexual intercourse in individuals experiencing fertility stress. In the study of Duman and Koçak (2013), it was determined that the most common sexual problem in infertile women was arousal at a rate of 20.5% and orgasmic disorder in 19.5% of women. In the study of Ergin et al. (2018), women reported that they did not find themselves attractive because sexuality was seen as a duty. In the process of infertility, couples who adapt to the change in sexual life increase mutual communication, attachment and satisfaction with marriage. It is stated that infertile women have lower marital adjustment than infertile men (Egelioglu Çetışli et al., 2014; Bodur et al., 2013) and that as dyadic adjustment increases in infertile couples, there is a decrease in anxiety and depression levels (Bodur et al., 2013).

Economic Problems in Infertile Couples

Since the drugs used in the treatment of infertility and the treatment process are quite expensive, they also cause economic difficulties for couples. Working couples are constantly dismissed from work due to treatment.

They have to get permission. Especially women quit their jobs or give up on making a career. In a study conducted by Koçyiğit in 2012, it was reported that the average expenditure for infertility treatments reached approximately three times the average household budget (Koçyiğit, 2012). The economic difficulties experienced can cause domestic economic violence. In a study, it was stated that 41 out of 122 infertile women were subjected to domestic violence and 29.2% were exposed to economic violence. In his study, Can (2005) found economic problems at a rate of 16.4% among the causes of stress experienced by infertile women during treatment. In the study of Kamacı (2003) it was determined that 84% of women and 85% of men stated that infertility treatment forced their family budgets economically.

Quality of Life of Infertile Couples

For most couples, infertility is a major life crisis. Studies on the relationship between the psychological reaction to infertility and gender have shown that women experience higher levels of anxiety, depression and loss of self-confidence compared to their spouses. However, psychological factors such as anxiety, depression and anxiety may affect physiological factors by reducing the probability of pregnancy. Psychosocial effects in the diagnosis and treatment of infertility can significantly affect the overall quality of life. In addition, spouses may react differently to infertility.

On the one hand, infertility itself, on the other hand, examination and treatment approaches applied for reproductive assistance, strain the coping skills and social support resources of the individual and the couple, and consume their physical and emotional energy, causing sexual dysfunction, depression, anxiety and deterioration in the couple's relationship. In the Rashidi and Montazeri study in Iran (2008), when the quality of life of women and men who receive IVF treatment is examined, according to the findings obtained, men lead a better and better quality life than women. In general, the relationship between being young, being female, low education level and quality of life was found to be significantly significant; No relationship was found between the duration of infertility, previous infertility treatment, or the causes of infertility and quality of life. It is understood that while the studies carried out to date mostly focus on documenting the physiological causes of infertility, less attention is paid to the individual's quality of life.

Johansson et al., examining the quality of life in couples 4–5.5 years after failed IVF treatment, found that childless couples had a lower quality of life than couples with children. Khayata et al. reported that female factor infertility adversely affects quality of life in infertile women compared to male and combined factor infertility.

Infertility Nursing

Nurses have an important role in supporting the communication of infertile couples, providing counseling for psychological problems, supporting coping resources and raising

awareness of the society. In addition to the roles of infertility nurses in medical care, there are roles of providing information, coordinator-manager, researcher, patient rights advocate, educator-counseling and psychological support. Within the scope of these roles, nurses often take responsibility in practices that require special knowledge and skills, such as evaluating the infertile couple, planning the determined treatment, assisting in assisted reproductive treatment procedures, preventing complications, case management, infertility counseling and coordinating. In addition, infertility nurses can provide information, explanation and decision-making counseling (the type of counseling in which the effect of starting or ending infertility treatment on the couple's life is explained), support counseling (the type of counseling given with therapeutic counseling for stress reduction) and therapeutic counseling (individual counseling, couple counseling). , group therapy, individual behavioral groups, emotional interactive groups, groups created using technology), telephone counseling and self-help groups are recommended. Nurses should be aware of the problems and effects caused by infertility, which creates a crisis in the lives of couples. The communication of the infertile couple should be supported, stress, anxiety and depression symptoms should be aware of the psychological problems that the couple may experience, risk factors should be questioned, and cooperation should be made with a psychologist or psychiatric nurse. The family should be supported in coping with the crisis situation, approaches that increase women's self-esteem and strengthen the relationship between couples should be adopted to reduce hopelessness, and public awareness should be raised in order to reduce social pressure on the couple.

Due to their changing and developing roles in the team, infertility nurses need to be aware of the economic and ethical problems and develop themselves in counseling for sexual health. Effectively fulfilling the roles of nurses will help couples to cope with problems and become stronger in coping with the process. Continuous trainings to be organized in this field will also contribute to the empowerment of infertility nurses.

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Quality of Life and Nursing Care in Menopause

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Introduction

Menopause, which is a natural part of the aging process in women, is defined as the loss of reproductive ability due to regression in ovarian functions. WHO defines menopause as “the permanent cessation of menstruation as a result of loss of ovarian activity”. In the studies, it is seen that the age of menopause varies between 49.3-51.4 years in developed countries, while this age range is 43.5-49.4 years in developing countries. When the TDHS data is examined, it has been determined that the age of Turkish women at menopause is concentrated in the 46-49 age range. According to the TDHS 2018 results, the age group with the highest menopause rate is the 48-49 age group with a rate of 45.1%. While the rate of women entering menopause in the 46-47 age range is 31.2%, the rate in the 44-45 age range is 15.9%.

In some women, the menopause process begins earlier than the expected age range. This process, called premature menopause; It is defined as “the loss of ovarian function of the ovary before the age of 40 (cessation of menstruation)”. According to the 2013 results of TDHS, the rate of women between the ages of 30-39 who entered menopause is 1.9%. According to the 2018 results of TDHS, the proportion of women between the ages of 30-39 who entered menopause increased to 2.6%.

As in every life period, many factors are effective on the time of menopause. In the fact that the age of menopause differs from society to society, even in different cultures of the same society; It is thought that individual factors such as genetic and environmental factors, nutrition and routine habits also have an effect. age of menopause; It is not affected by race, socio-economic status, number of pregnancies, oral contraceptive use, education, physical characteristics, alcohol consumption, age, or date of last pregnancy. It has been determined that smoking definitely accelerates the consumption of follicles and causes early menopause. Menopause directly affects both work and family life and social life of women.

Menopausal period; women experience intense hormonal changes; It is a period in which physiological, psychological, sexual and social changes occur depending on this change. The main reason for these changes is the decrease in estrogen level. During this period, symptoms related to different body systems may occur, from vasomotor symptoms to mental symptoms, from sexual function changes to osteoporosis. Symptoms such as menstrual irregularities, vaginal dryness, flora changes, sleep disorders, headache, dizziness, shortness of breath, heart palpitations, tremors, and anxiety can also be seen.

It is among the most important responsibilities of the nurse to determine their quality of life and the factors affecting it, while giving care to women in the menopausal period, and to use the results to improve women's health. Nurses should have knowledge about the risk factors that

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may develop by determining the changes that may occur in women in this period and know how this will affect their quality of life; will provide support for women to gain coping skills with menopausal complaints. In addition, nurses; Prioritizing the quality of life while giving care to women in the menopausal period will enable them to provide higher quality care and counseling services in order to increase the quality of life of women. In this section, the effects of menopausal problems on quality of life and nursing approach are discussed.

Changes Seen During The Menopause Period

With the menopause period, the activity of the ovaries begins to decrease and there is a decrease in the estrogen level to women; physiological, psychological, sexual and social etc. changes in many ways. These changes, which are thought to occur due to estrogen deficiency, cause many problems in the early and late periods. In the early period, vasomotor symptoms such as hot flashes, sweating, palpitations, headache, insomnia, muscle-bone pains, depression, loss of attention, forgetfulness, decreased libido, vaginal atrophy and urinary problems are. In the long term, there is an increase in the incidence of osteoporosis, cardiovascular diseases and cancers.

Cycle Changes

One of the most obvious differences experienced by women in the menopausal period is menstrual cycle changes. The termination of estrogen production, which is considered to be the main cause of menopausal changes, does not occur suddenly. Estrogen continues to be secreted in small amounts, so menstrual bleeding in women may continue for a while. This is most common in the premenopausal period. During this period, menstruation is of the anovulatory type. That is, the luteal phase has disappeared. Due to the decrease in the activity of the ovaries, enough estrogen is not secreted and the endometrial cycle cannot occur because the amount of estrogen in the blood decreases. Due to the anovulatory cycle, estrogen is constantly secreted from the follicles, and since this excess is not met by progesterone, bleeding can be intense and continuous during this period. As a result of all these changes, menstruation in women; It may occur as polymenorrhea, amonorrhea, hypomenorrhea, and hypermenorrhea. This type of bleeding can be a normal physiological process as well as a harbinger of various diseases. Therefore, women in the menopausal period should be considered as a risky group starting from the premenopausal period and the care to be given should be planned correctly.

Vasomotor Changes

Vasomotor changes are one of the most common symptoms, the severity of which varies by most women during the climacteric period. Vasomotor symptoms begin to appear from the premenopausal period and their incidence varies between 60% and 85%. vasomotor changes; Hot flashes are defined as an uncomfortable feeling of warmth that starts in the face, head or chest and progresses in any direction, sometimes spreading to the whole body. In the study of Ege et al. (2014) on women in the menopausal period; 69.5% experienced hot flashes, 55.3% had night sweats and 53.7% had insomnia problems; According to the study of Gözübebek and Başer (2016), it was seen that these vasomotor problems experienced by women negatively affect their daily lives.

Emotional Changes

Each life period has its own characteristics and stress factors. The menopausal period is also a period that includes many changes in women's lives. Therefore, it is important for women to be able to adapt to these changes and not to deteriorate their quality of life. The degree to which the woman is affected by each physiological event seen in this period and the meaning attributed to it varies from person to person. Mental problems are very common in the menopausal phase of women's life, especially in the perimenopause period. In the first place, thoughts that anxiety, depression and sexual dysfunctions increase are frequently expressed. Studies conducted to determine the emotional changes of women in the menopause period have also revealed that

women in this period experience emotional problems. In the study of Simon and Reape (2009), 91% of women in the climacteric period had irritability, 86% depression, 82% loss of concentration, 81% personality change, 77% sleep disorder, 77% It was determined that there was lack of motivation in .75%, memory deficit in 75%, hot flashes in 37%, headache in 19% and excessive sweating in 18%. According to the research results of Bozkurt and Sevil (2016); 44.2% of the women participating in the study had menopause after distress-stress, followed by “relaxation (20.8%)”, “inability to bear children (12.5%)”, “aging (8.3%)”, “a natural event (7.5%)” and “loss”. (6.7%)”. In Taiwan, a study was conducted with 3359 women aged 40-55 years to determine the level of depression occurring during the menopause, and 145 of the women had high-grade depression symptoms related to menopause.

Structural Changes in Bone

Another change that occurs as a result of long-term estrogen deficiency is osteoporosis. osteoporosis; It is a chronic and progressive musculoskeletal problem that begins with a decrease in the volume, density and quality of bone mass and continues with increased susceptibility to fracture. 50% of individuals aged 50 and over living in Turkey have osteopenia and 25% have osteoporosis. While this rate is 12.9% for women over 50 years old, it is 7.5% for men. Since 80% of people with osteoporosis are women, osteoporosis is an important gynecological problem. The national osteoporosis organization states that there are 200 million people in the world suffering from osteoporosis. When we look at the rate of women experiencing this problem, 30% of women with osteoporosis are postmenopausal women, and one out of every three women over the age of 50 experiences bone fractures caused by osteoporosis.

Cardiovascular Changes

Menopause is accepted as a risk factor for cardiovascular system diseases. Especially in terms of coronary artery disease, women with surgical menopause and early menopause are at higher risk than women who enter menopause in the normal age range. Since the positive effect of estrogen will disappear earlier in women who enter menopause early, these women are at higher risk for cardiovascular diseases. In the study of Ebong et al. (2014), the risk of heart failure in women with early menopause was found to be 66% higher than in women with normal menopause. Physiologically, estrogen decreases the LDL cholesterol level and increases the HDL cholesterol level, thus regulating the serum cholesterol level. In postmenopausal women, where estrogen levels are not sufficient, the cholesterol level gradually rises and the risk of atherosclerotic disease increases. Also, from menopause, women are more susceptible to weight gain and fat distribution due to changes in estrogen levels. This is another risk factor that puts women at risk for cardiovascular diseases.

Urogenital Changes

Another change that occurs due to a decrease in estrogen is urogenital changes. Urogenital changes; They are changes in the vulva, vagina, cervix, uterus, tuba uterine, pelvis and urinary system. The main reason behind these changes is the decrease in estrogen level. Among the most important effects of estrogen on the urogenital system of women; By increasing sexual motivation, it improves secondary sex characters and gives a feminine identity. Especially in women with early menopause, the absence of estrogen stimulation for a long time with menopause adversely affects the reproductive organs and regression is observed in the reproductive organs. This regression mostly occurs in the uterus, vagina, vulva and urethra. The vulva shows very rapid atrophy in the face of estrogen deficiency. Therefore, dystrophies are one of the most common problems in menopausal women in the future. Dystrophies cause excessive stenosis of the vulva and are accompanied by itching. In addition, atrophy seen in the vulva can cause cancer formation. With the decrease in estrogen level, some changes occur in the color, texture and flexibility of the vagina. These changes are more; It manifests itself as a narrowing in the vagina, a decrease in its color and

flexibility. In menopause, the vaginal wall becomes thinner and holds less moisture. This causes vaginal dryness. Vaginal dryness is seen at low rates in women with regular menstrual cycles; The incidence of vaginal dryness is increasing in perimenopause, postmenopause and late postmenopause periods. Dyspareunia is seen with vaginal dryness. In order to prevent dyspareunia, it is recommended to use water-soluble oils to eliminate vaginal dryness, and not to use insoluble oils such as vaseline. Because insoluble oils can clog the vaginal glands and cause infection. In addition, with the decrease of Estrogen, alkaline PH occurs in the vagina and the vaginal flora deteriorates. Estrogen deficiency changes the vaginal pH from 3.5-4.5 to 6-8. The alkaline environment makes the vagina susceptible to the invasion of a large number of pathogenic bacteria. This directly makes women more susceptible to infections. As a result of the loss of flexibility of the vagina after menopause, the infrastructure of the urethra shortens and narrows. Accordingly, especially in the following years, women; problems such as urination difficulties, true stress incontinence and urethral syndrome.

Vaginal disorders mentioned in postmenopausal women negatively affect sexual function. Levine et al. (2008). reported that sexually active women with sexual dysfunction and in the postmenopausal period were approximately four times more likely to have vulvovaginal symptoms than those without sexual dysfunction. In the same study, it was determined that 40% of women with vulvovaginal symptoms experienced general sexual dysfunction, 24% aversion, 34% difficulty in arousal, and 19% difficulty in orgasm [95]. In the study of Süt and Küçükkaya (2018), in which they investigated the sexual quality of life of women in the menopause period, it was determined that the most intense vaginal dryness complaint was in the perimenopausal period, and the complaint of dyspareunia was in the postmenopausal period, and it was reported that these symptoms reduced the quality of life of women [96]. Simon et al. (2014) in a study of 1000 postmenopausal women to investigate the effects of vulvovaginal atrophy, they reported that 64% of women experienced painful intercourse and loss of libido [89]. As can be seen, it is noteworthy that the quality of sexual life decreases in most of the women in the menopause period. Although these symptoms are not life-threatening, they can be progressive. All these urogenital system changes, in fact, seriously affect the sexual life of women and their quality of life. Women should develop some healthy lifestyle behaviors and evaluate treatment options in order to be least affected by these changes.

Dermatological Changes

One of the areas most affected by estrogen deficiency is the skin. Along with the decrease in estrogen level, the aging of the skin accelerates, collagen connective tissue and elastic fibers decrease, and the epidermis becomes thinner. Depending on the decreased collagen, epidermal changes such as decrease in the thickness of the epidermis, decrease in the secretion of the sebaceous glands, loss of flexibility, and decrease in blood flow are experienced. Studies show that HRT has a positive effect on skin dryness and skin thickness. It has been observed that there is a 10%-20% reduction in skin thinning and skin dryness in women taking HRT in the postmenopausal period (Aboobac. The decreasing estrogen level since menopause increases the testosterone level by decreasing the SHBG level, which leads to hair growth in the mustache and chin area and thinning of the hair in the pelvic and axillary region. Although all these skin changes are not actually a disease or a deterioration in physiological function, conditions such as thinning of the skin, wrinkling of the skin, deterioration in the appearance of the skin, sagging of the breasts, etc. can negatively affect the self-perspective in women, as they may cause a feeling of loss of attractiveness. and can damage self-esteem.

Menopause and Quality Of Life

It is very difficult to define quality of life and there is no agreed definition at present. The reason for this is that quality of life is a multidimensional concept used in all areas of life. According to the World Health Organization, health is “a state of complete physical, mental and social well-

being and not merely the absence of disease or infirmity”. Based on the “well-being” in this definition, WHO defines quality of life as “the patient's personal perception of his/her situation in life, both in the context of the cultural structure and value system in which he/she lives, and in terms of his/her own goals, expectations, standards and concerns”.

During menopause, women experience many symptoms and their daily living activities are significantly affected. Although the reasons for the emergence of menopausal symptoms are not known exactly, it is generally thought to be caused by the deficiency of ovarian hormones. These symptoms are; cycle disorders, vasomotor symptoms (VMS), emotional changes, sleep disorders, genitourinary problems, sexual problems, cardiovascular diseases and osteoporosis. All these changes bring about problems that negatively affect women's quality of life. Although the symptoms of menopause differ in each woman, it is estimated that 80-85% of women experience these symptoms and this reduces their quality of life. Physical, psychological and social changes that negatively affect the quality of life during menopause are quite complex. The nursing approach in this period includes careful diagnosis, support and education. Therefore, women's health nurses should provide care and counseling in order to increase the quality of life of women and to ensure that they spend this period in the best way possible. There are many studies in the literature to determine the symptoms and quality of life experienced by women during menopause. Ertem (2010) stated in his study that 79% of women in the menopause period experienced hot flashes and their quality of life decreased during this period. Asadi et al. (2012) found that the symptoms experienced by women during menopause were hot flashes (59.5%), mood changes (42.6%), vaginal dryness (41.1%), sleep problems (40.4%), night sweats (38.2%), memory loss (42.6%). 32.3%, urinary symptoms (18.3%), palpitations (6.6%), anxiety (5.8%), joint and muscle pain (59.9%), depression (4.4%), and irritability (3.6%). In the study conducted by Tümer and Kartal (2018) with 152 women to determine the relationship between women's attitudes towards menopause and their menopausal complaints, it was found that women experienced the most psychological problems. In another study, it was determined that 80% of women experience hot flashes and 31.3% of them experience night sweats. In an international study conducted in five European countries, it was reported that 55% of menopausal women and 75% of women living in the United States experienced hot flashes. In a study in many developed countries, it was reported that 75% of menopausal women experience hot flashes and 39% experience sweating. In the study conducted by Kharbouch and Hotun Şahin (2007), it was determined that women who have been in the postmenopausal period for five years or more have low quality of life scores in psychosocial, physical and sexual areas. All these study results show us that women may encounter many health problems during menopause, which is a normal process of their life stage, and their quality of life may be adversely affected.

Menopausal Period and Nursing Approach

Menopause is a period in which many important changes and related problems are experienced. The nurse, who is a member of the multidisciplinary team, helps women in the menopausal period to cope with these problems; practitioner, manager, trainer, consultant and researcher. Nurses, who are professional members of the health team, should play an active role in determining the health problems of women specific to the menopausal period, planning and implementing the interventions. Researches made; It has been shown that many women do not know what menopause is and what changes are taking place in their bodies. Based on the idea that many women have little knowledge about menopause, they should be told what changes will occur in menopause and how to take precautions, and whether these changes are normal.

It is among the important responsibilities of nurses to determine the quality of life of women in the menopausal period and the factors affecting them, and to use the results to improve women's health. The fact that nurses have knowledge about the quality of life of women in this period and are aware of the risks that affect their quality of life will enable them to provide more

support to women in coping with menopausal complaints. The nursing approach in this period includes careful diagnosis, support and education. Nurses should provide quality care and counseling services to improve the quality of life of menopausal women.

Health professionals should have up-to-date information about the menopausal period, menopausal symptoms, complications and their effects on quality of life. It is recommended to educate women on the management of symptoms related to menopause and how to prevent complications, to provide counseling to increase women's quality of life, and to carry out experimental studies that will lead to a personalized healthy lifestyle, considering that individual characteristics may affect menopausal complaints.

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The Relationship Between Long-Term Care Expenditures and Health Status

Gülay EKİNCİ¹

Introduction

Long-term care (health) consists of a set of medical and personal care services consumed for the purpose of relieving pain and suffering and managing deterioration in health status in patients with long-term dependence (OECD, 2011).

LTC includes services for patients with persistent or recurring illnesses and increased levels of addiction. These diseases include a wide range of services such as recurrent long or short-term care needs such as substance abuse, psychiatric disorders, management of chronic diseases such as cancer, end-of-life care, and care needs of individuals with physical disabilities.

Since long-term health services are provided informally (by family or relatives in the home environment) in low- and middle-income countries, data on these services are limited to upper-income countries where they were formal structured.

The need for long-term care is increasing day by day due to changes in demographic indicators such as the prolongation of life expectancy, the increase and structural changes in the burden of diseases, and the transformation in the work and employment structure.

The aging phenomenon, which develops due to the prolongation of life expectancy, increases the need for care support due to chronic diseases and mobility weakness due to aging. At the same time, the prevalence of chronic diseases such as high blood pressure, diabetes, muscle and bone diseases, alzheimer's, demenias increases the need for long-term care.

Long-term care services within the scope of "medical or nursing care" includes to prevent further deterioration of the patient's state of health, chronic disease management, rehabilitative care to restore functionality

Personal care services included supported self-care primarily due to disability and illness. These services are administered directly or under the supervision of nursing staff. These services provide help with activities of daily living (ADL) such as: eating, dressing, bathing, washing, getting in and out of bed, getting to and from the toilet and managing incontinence. These services are included within the health care boundary because the purpose of this type of care is linked to survival and the maintenance of health status (OECD, 2011).

Health status is an indicator of an individual's health level. Health status can be evaluated objectively and subjectively. The concept of perceived health status is a subjective concept.

In the perceived health status, the individual may have a positive or negative attitude towards the disease they have, or they may remain indifferent. Therefore, there are three types of

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assessments on perceived health status. It is possible for an individual to feel good, to feel bad, or to evaluate himself as neither good nor bad.

In this context, three types of evaluations are made under the heading of the individual's perceived health status in the OECD database. These perceived health status were "Good/very good health, total aged 15+ % of population (crude rate) "; "Fair (not good, not bad) health, total aged 15+ % of population (crude rate)"; "Bad/very bad health, total aged 15+ % of population (crude rate)".

Health status can be measured statistically and objectively expressed with various indicators and methods developed (Goodacre vd., 2010) These indicators are life expectancy, mortality rates, morbidity rates.

The need for long-term care is closely related to life expectancy and morbidity from a health status perspective.

Average Life Expectancy is one of the most common measures used to obtain information about the health of a population. Average life expectancy is a rough measure of how long a person will live. This measure shows how long a person is likely to live at a given age unless there is a trend change in death rates (Ulutürk, 2015).

Morbidity is another measure of the health status of the population. It gives the ratio of individuals with impaired health to the population of the society. The most important concept used within the scope of morbidity is disease burden. The burden of disease measures the difference between the current life status and the ideal life status, where each individual can live free from diseases and disabilities in later life, to see the effect of diseases and disabilities (Ulutürk, 2015). The most frequently used indicators of disease burden from the perspective of health economics;

- Quality Adjusted Life Year (QALY) is defined as the expected life expectancy of the patient and the quality of life in these years. QALY value is between 1 and 0. "1" represents perfect health while "0" represents death (Rudmik ve Drummond, 2013). For example, if a heart disease that has a negative impact on quality of life reaches a disability score of 0.8, the quality of life score will be 0.2.
- Disability Adjusted Life Year (DALY) is a measure of the overall disease burden, expressed as the number of years lost due to loss of health, disability, or premature death.
- Year Lost Life (YLL) expresses the number of deaths occurring at each age multiplied by the estimated remaining life years according to life expectancy. It is based on the calculation of life years lost due to death.
- Year Lost Disability (YLD); the number of cases resulting from injury or illness is based on disability-adjusted life years calculation.

Diseases are considered at three basic levels in the calculation of disease burdens:

a. Noncommunicable diseases (NCD) are a group of noncommunicable diseases that occur in the long term due to one or more risk factors, for example; Alzheimer's, diabetes, cancer diseases etc.

b. Communicable diseases—CD is a type of disease that has a disease agent in its etiology, occurs in the short or long term, and shows contagious features, for example; AIDS, syphilis, tuberculosis, influenza etc.

c. Injuries are the negative changes that occur in the health level of the individual by any sharp object injury, fall, drowning, poisoning, accident, terrorism, etc. For example; traffic accidents, drowning, etc.

At a global level, DALY for non-communicable diseases was 1.6 billion/year, DALY for communicable diseases was 668 million/year, and DALY for injuries was 249 million years was calculated. DALY of non-communicable diseases has a high share of 64% in the total disease burden. This calculation shows that the years of life spent due to diseases related to chronic diseases or lost due to premature death are at a very high level. Therefore, chronic diseases appear as an issue that should be evaluated in terms of health economics.

According to the World Health Organization (WHO), chronic diseases are defined as diseases that are not contagious, persist for 6 months or more, generally progress slowly, and many risk factors play a role in their development.

According to the Commission on Chronic Illness, chronic diseases are defined that are generally non-infectious in nature, in which socioeconomic, personal characteristics, and genetic factors play a role in the formation of slowly progressive, permanent, and usually permanent disabilities, in which the recovery is not usually seen completely (Aywater, 1949; Bilir, 2006).

The group of non-communicable diseases consists of chronic diseases. Among these diseases, mental diseases such as cancer, cardiovascular system diseases, diabetes, Alzheimer's, and dementia have an important place. Chronic diseases involve high costs due to the long-term diagnosis, treatment, and rehabilitation processes. In addition, increases in life expectancy and the incidence-prevalence of chronic diseases brought about by these increases also increase health expenditures.

In a study, the annual cost per patient was calculated as \$48293 for HIV/AIDS disease; \$38551 for kidney diseases, \$37155 for CVD, and \$25016 for respiratory disorders (Hajat et al, 2021).

These diseases involve high costs depending on their duration and whether they are comorbid with additional diseases.

In the same study, in the evaluation of chronic diseases per patient; the Annual health expenditure of patients with 2 chronic diseases was 4385 dollars, 5851 dollars in 3 chronic diseases, 10088 dollars in 5 chronic diseases, 19856 dollars in 8 chronic diseases, 33874 dollars in 11 chronic diseases (Hajat et al, 2021).

In another study, in the percentage of Health Care Total Spending by Number of Chronic Conditions, it was determined that; 16.2% of expenditures are spent on 0 chronic diseases, while 18.4% on 1 chronic disease; 17.2 % was spent on 2 chronic diseases, 15 % on 3 chronic diseases, 11.1 % on 4 chronic diseases, and 22.1 on 5 or more (Gerteis et al. 2014).

Countries make health expenditures to manage the health needs for individuals. Health expenditures are generally considered as total and current health expenditures. However, demographic changes and changing health needs cause the services offered to diversify or to be handled in detail. Contrary to the general approach to health expenditures, WHO has published a more detailed guide and discussed health expenditures in 9 categories. One of these expenditures is long-term maintenance expenditures. Long-term care expenditures are the expenditures made to ensure the well-being of the individual for conditions such as the management of chronic diseases, the need for palliative care services, age-related health deficiencies, and substance abuse.

In this context, long-term care expenditures are handled in two dimensions health expenditures and social expenditures. Long-term care expenditures-social are expenditures made to meet the social and economic needs of individuals (retirement, incapacity payments, sickness payments, services to support daily activities, etc.) in long-term deterioration in the health status of the individual. Long-term care expenditures-health are health-related expenditures made for the management of long-term deterioration in the health status of the individual (OECD, 2011). Long-

term care-health; i) inpatient long-term care-health, ii) Day long-term care-health, iii) Outpatient longterm care-health, iv) Home-based longterm care-health.

Countries determine their health expenditures and investments by taking into account the health services needed by the population. This study, it was aimed to evaluate long-term care expenditures, life expectancy at birth, and perceived health status.

Methods

Descriptive methods were used in the analysis. The analysis was done in three stages. In the first stage, descriptive information about the variables was given; in the second stage, the increase-decrease relationship of variables was examined according to years; in the third stage, the correlation relationship between the variables was examined; in addition, the relationships between the variables were evaluated through graphics.

Variables

Only 29 countries had regular long-term care expenditures data between 2010-2020. These countries were Austria, Belgium, Canada, Croatia, Czech Republic, Denmark, Estonia, Finland, France, Germany, Greece, Hungary, Iceland, Ireland, Korea, Latvia, Lithuania, Luxemburg, Netherlands, Norway, Poland, Portugal, Romania, Slovak Republic, Slovenia, Spain, Sweden, Switzerland, United States. The data on long-term care expenditures according to the PPP\$ (Per capita, current prices, current PPPs) for 2010-2020 was used.

The variables used in the study were long-term care expenditures, life expectancy at birth, three separate data given under the heading of Perceived health status as the well-being of the population in terms of morbidity. These variables of Perceived health status were "Good/very good health, total aged 15+ % of population (crude rate) "; "Fair (not good, not bad) health, total aged 15+ % of population (crude rate)"; "Bad/very bad health, total aged 15+ % of population (crude rate)".

Limitations Of This Study

The limitation of this study was considered to be the number of countries and year range. The other limitation of this study is Turkey does not have data on long-term care expenditures, so it could not be included in the analysis. In addition, disease burdens were not evaluated as a variable in this study.

Statistical Analysis

Statistical analyzes were made using Eviews 10 program (Eviews 10, IHS Global Inc., 4521 Campus Drive, #336, Irvine, CA 92612).

Results

Results were made in three stages such as; i) descriptive analysis of the variables, ii) results on the relationship between long-term care expenditures and variables , iii) results on the relationship between life expentancy at birth and variables.

a. Descriptive analysis of the variables

According to Table 1; the mean of LEAB of 29 countries was 80.01 ± 2.71 (min: 73.10; max: 84.00); the mean of LTCE was 595.26 ± 494.08 PPP\$ (min: 6.42; max: 1946.49); the mean of Good Health Status was 67.02 ± 13.11 (min: 29.50; max: 89.00); the mean of Not Good Not Bad Health Status was 23.16 ± 8.86 (min: 8.00; max: 53.50); the mean of Bad Health Status was 9.86 ± 4.96 (min: 2.50; max: 26.90).

Tablo 1. Descriptive Analysis of The Variables

	LTCE*	LEAB	Good Health	Not Good Not Bad	Bad Health
Mean	595.2613	80.01379	67.01661	23.16426	9.858934
Median	636.3710	81.10000	69.50000	21.70000	8.700000
Maximum	1946.494	84.00000	89.00000	53.50000	26.90000
Minimum	6.423000	73.10000	29.50000	8.000000	2.500000
Std. Dev.	494.0802	2.706282	13.10574	8.864702	4.964747
Skewness	0.487303	-0.802500	-0.637659	0.964884	0.692728
Kurtosis	2.168927	2.449334	3.000417	4.239605	3.111243
Jarque-Bera	21.80549	38.27010	21.61803	69.92253	25.67768
Probability	0.000018	0.000000	0.000020	0.000000	0.000003

Source: Prepared by the author, *Per capita, current prices, current PPPs

Long-term care expenditures have increased by an average of 1.02 times (102%) at the level of 29 countries in 11 years. Countries with the highest increase in Long-Term Care Expenditures in 2020 compared to 2010 were Croatia; Czech, Estonia, Korea, Greece, Poland, Germany (an increase of 1x or more); decreases were experienced in Luxemburg and Romania.

Life Expectancy At Birth has increased by an average of 1.3% at the level of 29 countries in 11 years. Countries with the highest increase in Life Expectancy At Birth in 2020 compared to 2010; Korea, Estonia, Latvia, Denmark, Finland, Lithuania, Norway, Ireland (an increase of 2% or more); decreases were experienced in Romania and United States (respectively %0.27; %2.1)

Good Health Status has increased by an average of 2.7 % at the level of 29 countries in 11 years. Countries with the highest increase in Good/Very Good Health total aged 15+ in 2020 compared to 2010; Croatia, Hungary, Slovenia, Estonia (an increase of 10-30 %); the highest decreases were experienced in Korea, Lithuania (decreases respectively 16%, 12%).

Fair Health Status has increased by an average of 1.08 % at the level of 29 countries in 11 years. Countries with the highest increase in Fair (Not Good/Not Bad) Health total aged 15+ in 2020 compared to 2010; United States, Luxemburg, Sweden, Korea, Lithuania, Portugal (an increase of 10-30 %); the highest decreases were experienced in Slovenia, Austria, Switzerland, Ireland, Germany (decreases respectively 15-10 %).

Bad Health Status has decreased by an average of 10.7 % at the level of 29 countries in 11 years. Countries with the highest increase in Bad/very bad Health total aged 15+ in 2020 compared to 2010; Germany, Ireland, Norway, Iceland, United States, Switzerland, (an increase of 2.8-52 %); the highest decreases were experienced in Croatia, Hungary, Portugal, Greece, Poland, Slovenia, Romania, Czech, Estonia, Finland, Latvia, Austria, Spain, Lithuania, Belgium, Netherlands, Denmark, Korea, Slovak Republic, Sweden, Canada, Luxemburg (decreases respectively 2.5-44 %); besides this there was no increase or decrease in France.

Tablo 2. Descriptive Analysis of The Series

Countries	LTCE	LEAB	Good/very good health, total aged 15+	Fair (not good, not bad) health, total aged 15+	Bad/very bad health, total aged 15+
Germany	1,0276	0,0075	-0,0215	-0,1049	0,5185
Ireland	0,3329	0,0223	0,0109	-0,1143	0,2188
Norway	0,4298	0,0259	-0,0287	0,0581	0,1538
Iceland	0,4689	0,0147	-0,0154	0,0353	0,1346
United States	0,4176	-0,0216	0,0034	0,1847	0,1000
Switzerland	0,4480	0,0048	0,0221	-0,1267	0,0286
France	0,5491	0,0049	0,0178	-0,0504	0,0000
Luxemburg	-0,0605	0,0173	-0,0239	0,1131	-0,0250
Canada	0,4965	0,0037	0,0102	-0,0889	-0,0345
Sweden	0,2520	0,0098	-0,0242	0,1235	-0,0370
Slovak Republic	0,4036	0,0185	0,0252	-0,0398	-0,0435
Korea	2,3092	0,0411	-0,1622	0,1584	-0,0495
Denmark	0,2811	0,0290	0,0028	0,0095	-0,0506
Netherlands	0,5585	0,0049	0,0000	0,0353	-0,1000
Belgium	0,4204	0,0062	0,0329	-0,0615	-0,1429
Lithuania	0,8456	0,0246	-0,1175	0,2990	-0,1818
Spain	0,2752	0,0000	0,0167	0,0149	-0,1852
Austria	0,3803	0,0074	0,0647	-0,1315	-0,1935
Latvia	0,8054	0,0328	0,0397	0,0548	-0,2171
Finland	0,1248	0,0224	0,0220	0,0041	-0,2222
Estonia	2,9224	0,0382	0,1082	-0,0679	-0,2349
Czech	5,3974	0,0077	0,0193	0,0664	-0,2377
Romania	-0,1314	-0,0027	0,0641	-0,0963	-0,2396
Slovenia	0,6274	0,0100	0,1258	-0,1365	-0,2857
Poland	1,1378	0,0013	0,0657	0,0299	-0,2987
Greece	1,4271	0,0099	0,0384	0,0068	-0,2990
Portugal	0,9169	0,0125	0,0407	0,1639	-0,3448
Hungary	0,4292	0,0134	0,1273	-0,0106	-0,4036
Croatia	6,1942	0,0143	0,3181	-0,0143	-0,4424
	1,0237	0,0131	0,0270	0,0108	-0,1074
*Year ranges used in the calculation of countries data: LTCE: Ireland 2011-2020, Croatia 2011-2020; Good Health Status: Iceland 2010-2018, United States 2010-2019; Fair Health Status: Iceland 2010-2018, United States 2010-2019; Bad Health Status: Iceland 2010-2018, United States 2010-2019.					

*Source: Prepared by the author***b. Results on the relationship between long-term care expenditures and variables**

The correlation analysis results of the variables were given in Table 3.

Table 3: Correlation Analysis Results

	LTCE	DBYS	GOOD VERY GOOD HEALTH TOTAL AGED 15	FAIR NOT GOOD NOT BAD HEALTH TOTAL AGED 15	BAD VERY BAD HEALTH TOTAL AGED 15
LTCE	1	0.64	0.62	-0.51	-0.68
DBYS	0.64	1	0.47	-0.38	-0.56
GOOD VERY GOOD HEALTH TOTAL AGED 15	0.62	0.47	1	-0.96	-0.90
FAIR NOT GOOD NOT BAD HEALTH TOTAL AGED 15	-0.51	-0.38	-0.96	1	0.76
BAD VERY BAD HEALTH TOTAL AGED 15	-0.68	-0.56	-0.90	0.76	1

Source: Prepared by the author

According to Table 3, there was a positive correlation between long-term care expenditures and life expectancy at birth ($r=0.64$). It is understood from Figure 1 that long-term care expenditures between 0-500 PPP dollars were concentrated in the 73 years and over, while the expenditures of 500 dollars and above were mostly made in the 78-84 age group (Figure 1).

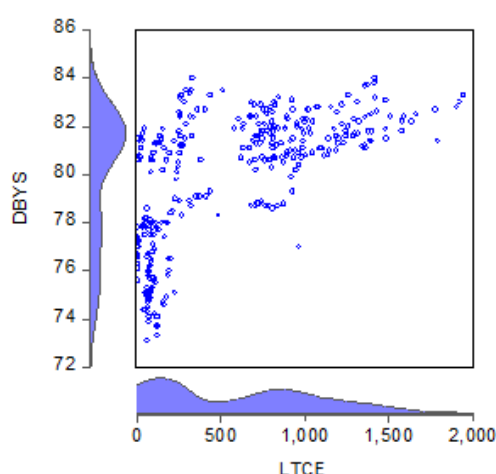


Figure 1. Long-term care expenditures and life expectancy at birth, 29 countries, 2010-2020

There was a positive correlation between long-term care expenditures and Good Health Status ($r=0.62$). According to Figure 2, while long-term care expenditures between 0-500 PPP dollars were pointed to the Good Health Status of 29.5-89 % population; spending 500 PPP dollars or more was pointed to the Good Health Status of 60-90 % population.

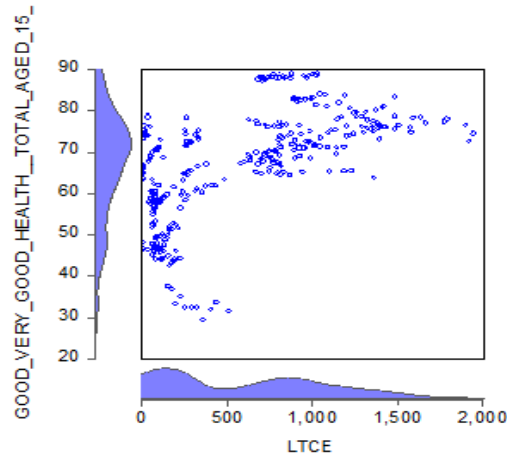


Figure 2. Long-term care expenditures and Good Health Total Aged 15+, 29 countries, 2010-2020

There was a negative correlation between long-term care expenditures and Fair Health Status ($r = -0.51$). According to Figure 3, while long-term care expenditures between 0-500 PPP dollars were pointed to the Fair Health Status of 15-55 % population approximately; spending 500 PPP dollars or more was pointed to the Fair Health Status of 8-30 % population approximately.

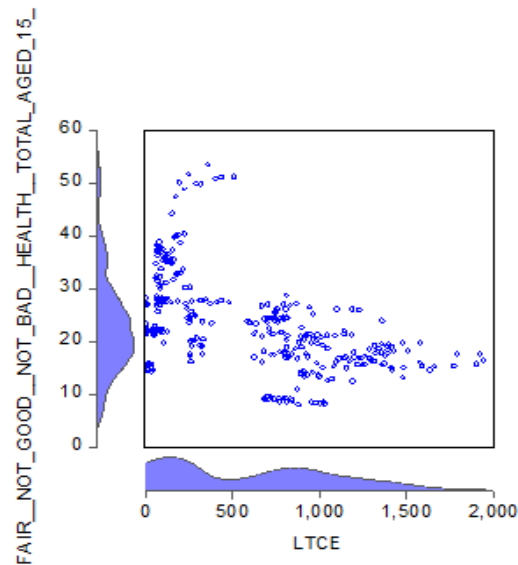


Figure 3. Long-term care expenditures and Fair Health Total Aged 15+, 29 countries, 2010-2020

There was a negative correlation between long-term care expenditures and Bad Health Status ($r = -0.68$). According to Figure 4, while long-term care expenditures between 0-500 PPP dollars were pointed to the Bad Health Status of 5-27 % population approximately; spending 500 PPP dollars or more was pointed to the Bad Health Status of 2.9-12 % population approximately.

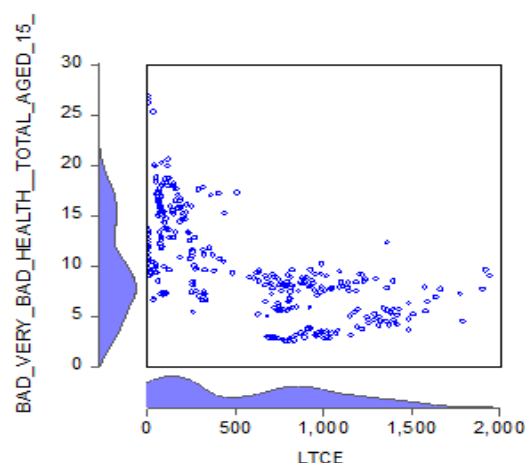


Figure 4. Long-term care expenditures and Bad Health Total Aged 15+, 29 countries, 2010-2020

c. Results on the relationship between life expectancy at birth and variables

There was a positive correlation between life expectancy at birth and Good Health Status ($r=0.47$). From Figure 5, it is understood that in the age range 80-84 around 60-90% of the population indicated Good Health Status.

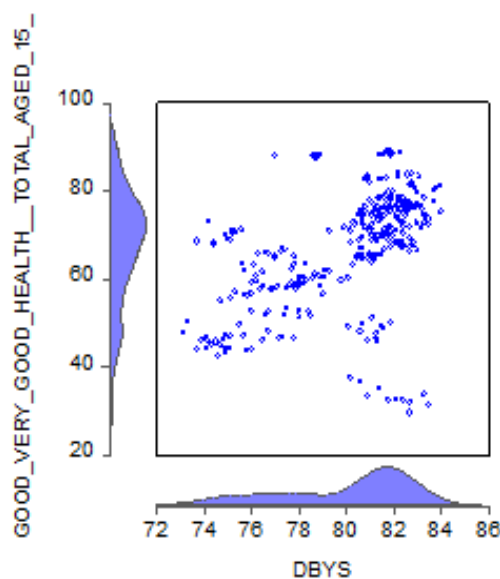


Figure 5. Life Expentancy At Birth and Good Health Total Aged 15+, 29 countries, 2010-2020

There was a negative correlation between life expectancy at birth and Fair Health Status ($r=-0.38$). From Figure 6, it is understood that in the age range 80-84 around 8-30 % of the population indicated Fair Health Status.

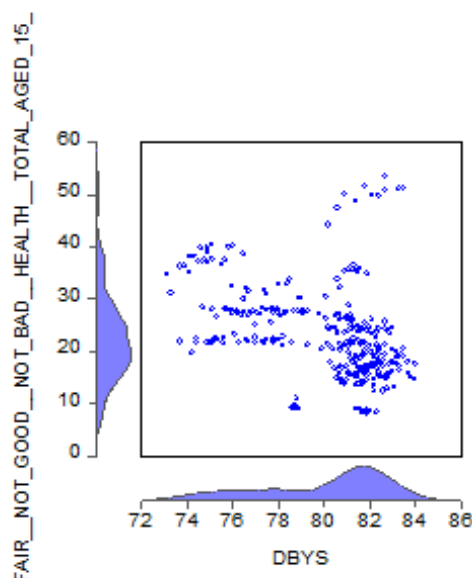


Figure 6. Life Expentancy At Birth and Fair Health Total Aged 15+, 29 countries, 2010-2020

There was a negative correlation between life expectancy at birth and Bad Health Status ($r=-0.56$). From Figure 7, it is understood that in the age range 80-84 around 2.5-22 % of the population indicated Bad Health Status.

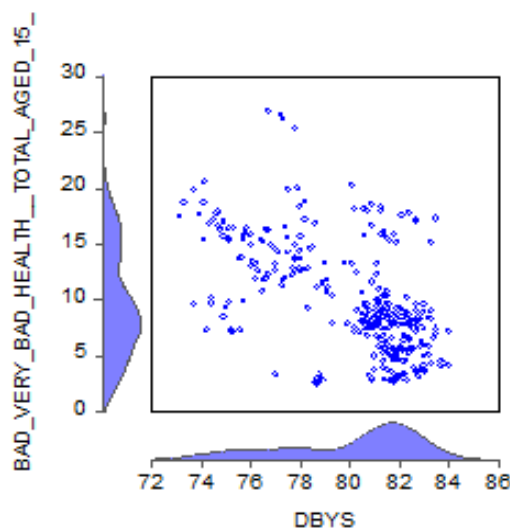


Figure 7. Life Expentancy At Birth and Bad Health Total Aged 15+, 29 countries, 2010-2020

Discussion and Recommendations

There are many influential factors on perceived health status such as age, gender, education, marital status, working conditions, genetics, diseases burdens etc.

In a study, when averages in health status perceptions are detailed according to rural-urban, gender, marital status, income, and education level, the most important factors affecting positive health perceptions are marital status, education level, and income level, respectively. In a study conducted in Turkey, the European Union, and OECD countries, approximately 30-32% of people reported that they had health problems, while 65-70% of the population stated that their health status was "good" (Aydın, 2019). In this study, our results showed that while 67% of the population perceived their health status as good in the countries subject to the study, 23.2% evaluated it as fair health status and 9.2% as bad health status.

Population satisfaction with regard to coverage of primary health care services at the level of 10 industrialized countries, 12 health indicators (eg infant mortality, life expectancy, and age-adjusted mortality rates) and overall costs of systems was studied. While the results reveal overall concordance for primary care, health indicators and satisfaction-expenditure ratio in nine out of 10 countries; USA and West Germany were found to be at low levels (Starfield, 1991).

In a study using Health Expenditure, Maternal Mortality, and Neonatal Mortality, it was concluded that health expenditures can improve the aforementioned indicators of low HDI countries by mediating the relationship between HDI and neonatal and maternal mortality rates (Nuhu et al, 2018).

In a study using Disability Rate, Mortality Rate, and Quality of Life, it was concluded that there is a strong relationship between Human Development Index and health indicators (Prashanth, 2021).

In the evaluation of 29 countries between 2010-2020;

- The average life expectancy in the countries studied is 80 years old.
- Long-term care expenses average around 600 PPP dollars.
- Long-term care expenditures have increased by over 100% in 11 years.
- Average life expectancy increased by around 1.3%.
- The percentage of the population with good health increased by around 2.7%.
- The percentage of the population with a fair health status increased by 1.08%.
- The percentage of the population with bad health status decreased by around 10.7%.

A moderate and positive correlation was found between long-term care expenditures and average life expectancy. Results showed that long-term care expenditures between 0-500 PPP dollars were concentrated in the 73 years and over, while the expenditures of 500 PPP dollars and above were mostly made in the 78-84 age group (Figure 1).

A moderate and positive correlation was found between long-term care expenditures and the percentage of the population with good Health status. According to Figure 2, while long-term care expenditures between 0-500 PPP dollars were pointed to the Good Health Status of 29.5-89 % population; spending 500 PPP dollars or more was pointed to the Good Health Status of 60-90 % population.

A low level and negative correlation was found between long-term care expenditures and the percentage of the population with moderate and poor health. In other words, while the increase in long-term care expenditures causes a decrease in the fair-bad health level; the decrease in long-term care expenditures causes an increase in the fair-bad health level.

A low level and positive correlation was found between average life expectancy and the percentage of the population with good health status. From Figure 5, it is understood that in the age range 80-84 around 60-90% of the population indicated Good Health Status.

A low level and negative correlation was found between average life expectancy and the percentage of the population with fair-bad health status. From Figure 6, it is understood that in

the age range 80-84 around 10-30 % of the population indicated fair health status. From Figure 7, it is understood that in the age range 80-84 around 2.5-22 % of the population indicated bad health status.

The results of the evaluation of the countries within the framework of the variables;

- Countries where long-term care expenditures and life expectancy at birth increase together, with a decrease in the level of good and fair health status and an increase in bad health status; Germany.
- Countries where long-term care expenditures and life expectancy at birth increase together, with a decrease in the level of good health status and an increase in the level of fair and bad health status; Iceland, and Norway.
- Countries where a decrease in long-term care expenditures and an increase in life expectancy at birth, with a decrease in the level of good and fair health status and an increase in bad health status; Luxemburg.
- Countries where an increase in long-term care expenditures and a decrease in life expectancy at birth, with an increase in good-fair-bad health status together; United States.
- Countries where long-term care expenditures and in life expectancy at birth increase together, with an increase in the level of good health status and a decrease in fair-bad health status; Croatia, Hungary, Slovenia, Estonia, Austria, Belgium, Slovak Republic, and Canada.
- Countries where long-term care expenditures and life expectancy at birth increase together, with an increase in the level of good-fair health status and a decrease in bad health status; Portugal, Greece, Poland, Czech, Denmark, Netherlands, Spain, Latvia, Finland.
- Countries where long-term care expenditures and life expectancy at birth increase together, with an increase in the level of good-bad health status and a decrease in fair health status; France, Ireland, and Switzerland.
- Countries where long-term care expenditures and life expectancy at birth increase together, with an increase in fair health status and a decrease in good-bad health status; Lithuania, Korea and Sweden.
- Countries where long-term care expenditures and life expectancy at birth decrease together, with an increase in good health status and a decrease in fair-bad health status; Romania.

Generally the increase in long-term care expenditures with the increase in life expectancy points to the percentage of the population with a better good health status. Therefore, it is thought that it will guide the policies to be followed for long-term care expenditures.

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Polycyclic Aromatic Hydrocarbons and Their Effects on Health

Hakan TOĞUŞ

1. Definition

Polycyclic aromatic hydrocarbons (PAHs) are structures formed by the combination of two, three or more aromatic rings. Under the influence of high temperature, pyrolysis of hydrocarbons occurs. PAHs with less than four benzene rings are defined as mild PAHs, and PAHs with four or more benzene rings are defined as severe PAHs (Wu, Tao & Liu, 2006). Coal, oil, cigarette or hookah smoke, smoke from industrial facilities, and foods cooked on barbecue are among the sources of PAHs (Li et al, 2006).

Polycyclic aromatic hydrocarbons are a semi-volatile group of organic pollutants that can be formed and spread through various agricultural activities and industrial areas (Wick et al, 2011). It has been found that it can easily contaminate the air, water, soil and nutrients (Srogi, 2007). PAHs are compounds with higher affinity for heavy metals and other organic pollutants due to their strong hydrophobic properties. Due to their low water solubility, they are difficult to detect, so accurate and appropriate analysis methods must be developed (Hu et al, 2015). PAHs can be formed in foods as a result of heat treatments such as barbecue, grilling, smoking and roasting, as well as the compounds found in polluted air conditions in industrial production areas, accumulating on plants and contaminating vegetables, fruits and cereals (Moazzen et al, 2015).

These groups of compounds have potentially harmful effects on ecosystems and human health, as many of them have been shown to be carcinogenic, teratogenic and mutagenic. (Sojinu, Sonibare & Zeng, 2011). Various approaches and strategies, including physical and chemical, have been developed, optimized and used to reduce the effects of these pollutants and rehabilitate polluted areas (Moazzen et al, 2015). Some of these traditional remediation techniques are not common due to their technological complexity, high cost, and significant limitations, such as lack of public acceptance. Most of the techniques are invasive and simply relocate the contamination problem, which often requires more waste management (Sojinu, Sonibare & Zeng, 2011).

2. History

Although the fact that chemicals such as PAH can cause cancer was determined as a result of a research conducted in London in 1775, there is not enough information about the damage of these compounds on the tissue. In 1920, it was determined that tumors formed on the skin of mice after soot extracts were applied to the skin of mice by Japanese workers. In 1953, it was found by Doll that smoking is the main cause of lung cancer. After the analysis, BaP in cigarette smoke and tar was found to have a carcinogenic effect (Simko, 2002).

3. Physical and Chemical Properties

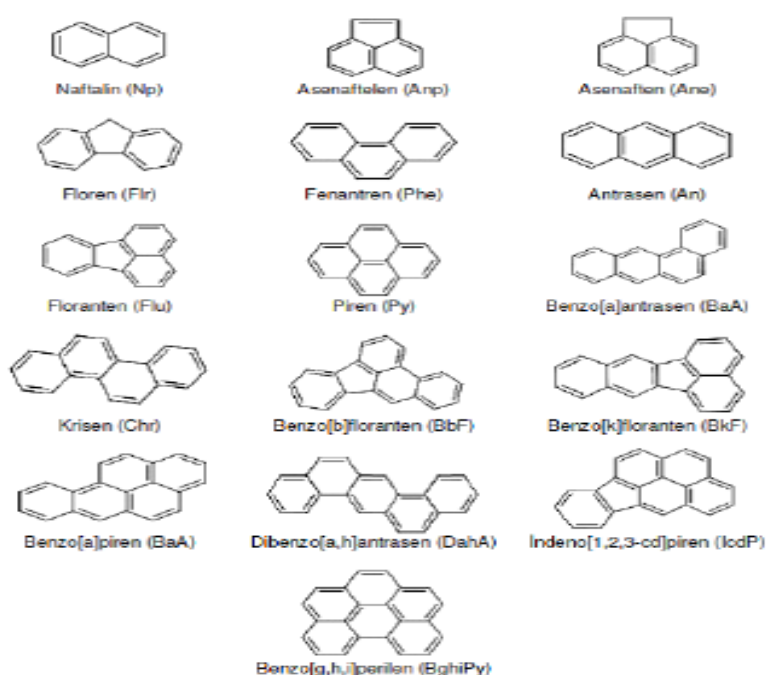
PAHs are compounds that occur as a result of incomplete combustion of hydrocarbons, and often occur as a complex mixture of combustion products instead of a single compound (ATSDR, 1995). But it can also be produced as a pure compound for research purposes. PAHs in pure compound form are white, green, light yellow, colorless, solid and have a slight pleasant odor. Most of these hydrocarbons do not have a specific area of use, except for PAHs produced for purposes

other than research. Some PAH compounds can be used in healthcare and in the manufacture of paints, pesticides or plastics (ATSDR, 1995). Figure 1 shows the molecular structures of some PAH groups. PAHs react with nitric acid and nitrous oxide to form sulfur dioxides, nitro derivatives, and sulphonic and sulfuric acid forms by reacting with sulfuric acid (WHO, 1998). In addition, they can react with ozone and hydroxyl radicals (Douben, 2003). Factors such as the solubility of PAHs in water, their diffusion in the environment, their vapor pressure, and logKow (WHO, 1998).

4. Classification

It is known that there are more than 100 PAH compounds in nature. However, 16 different PAH compounds in Figure 1, which are thought to have higher carcinogenic and toxic effects, are recognized as priority pollutants by the United States Environmental Protection Agency (US-EPA) (Rubailo et al, 2008).

Figure 1. 16 different PAH compounds thought to be carcinogenic.



5. Presence in Foods

PAH compounds formed due to environmental pollution (Szterk, 2013) in industrial production areas pass into the soil, water and air and contaminate agricultural products such as vegetables, fruits and cereals (Shi, Zhang & Liu, 2016). In addition, these compounds have been found in processed and heat-treated foods such as roasted coffee, dairy products, vegetable oil, tea, and packaging materials (Londoño, Reynoso & Resnik, 2015). PAH compounds formed in coffee occur during roasting of coffee beans (Szterk, 2013). PAH compounds found in oilseeds; As a result of contamination with environmental factors such as soil, water and air, it can infect these seeds, and it can be transferred to vegetable oils in stages such as drying method and solvent extraction during the seed stage (Shi, Zhang & Liu, 2016). European Commission regulations (835/2011); propolis etc. reported that various herbal products such as green tea and bee products contain PAH compounds depending on the production techniques (EC, 2011).

It is thought that PAH compounds can be formed in foods by basically three different methods. First method; It is the formation of PAH compounds as a result of pyrolysis of organic substances such as carbohydrates, proteins, fats, etc. with high temperature (500-900 °C). Second method; The oil droplets dripping from the foods cooked with coal fire can turn into volatile PAHs as a result

of dripping on hot coal and burning with intense fire. These volatile PAH products are more likely to contaminate the surface of the food as the smoke point increases. The third method is; The formation of PAH products with the incomplete combustion of coal and the contamination of these products on the food surface (Esposito et al, 2015). In particular, fats melted in meat cooked on barbecue or charcoal fall on hot coal and become pyrolysis (Alomirah et al, 2011). Thus, the PAH compounds formed are carried towards the meat by the smoke and accumulate on the surface. Later, it is carried into the food and accumulates in lipid components (Santos, Gomes & Roseiro, 2011). Due to the lipophilic nature of PAH compounds in this transport, the fat and water content of the food has an important role in the transport rate (Santos, Gomes & Roseiro, 2011).

Polycyclic aromatic hydrocarbons are environmental pollutants formed as a result of the effects of solid fuels, fossil fuels and biological organisms (Huang and Penning, 2014). In addition, smoking, cooking, stove use, and incense are also shown to be among the main sources of PAH (21). For this reason, the PAH rate is high in kitchens, and it can also be taken into the body from air, water, soil, food and water consumption (WHO, 2014).

6. PAHs in the Food Chain

The sources of PAHs in foods by contact with foods are numerous. PAHs are industrially released into nature as pyrolysis from organic substrate compounds originating from human activities. PAHs emitted to the environment as a result of the combustion of wood or coal and the emission of exhaust fumes from motor vehicles become contaminated with food (WHO, 2014).

Many PAH compounds have low water solubility and low volatility. Therefore, they tend to remain in the form of small organic particles such as ash. As a result of the contamination of these compounds with soil, foods such as legumes, cereals, vegetables and fruits are grown in contaminated soil (WHO, 2014). In a study conducted in England, the amount of B[a]P in the peel of vegetables as a result of soil contamination was found to be between 590 and 2301 µg/kg. The high amount of PAH in the soil causes contamination by passing from the plants that animals consume while grazing to their milk and from there to dairy products (WHO, 2014).

7. PAH Formation During Food Preparation and Cooking

Foods may be contaminated with PAH before preparation, but the PAH content in foods can often increase during cooking (Wick et al, 2011). In the researches on the formation of PAH in the cooking stages of foods; While the method that produces the most PAH is seen as roasting and frying, the least PAH formation is observed in the boiling method (WHO, 2014). The contribution of these methods to the PAH content can be explained as follows.

7.1. Food Smoking

The smoking method, which is one of the oldest cooking methods, is often used in the storage and flavoring of foods such as meat, fish and cheese. In this method, it is applied at 650-700 °C in wood fire. The smoke from the fire comes into contact with the food, and the food continues to be smoked as the temperature of the fire decreases (WHO, 2014). Liquid smoke flavorings (LSF) can also be used to reduce PAH formation in smoking. This method provides lower PAH formation compared to smoking (WHO, 2014).

7.2. Grilling

In this method, in which heat is transmitted to the food by using an oven or pan, high-temperature butter, margarine or vegetable oils can be used. During this process, PAH formation can occur mostly as a result of cooking fatty beef with barbecue (WHO, 2014).

7.3. Frying

The method of cooking foods with animal or vegetable oil at high temperature is called frying. In this cooking method, since the heat in the hot oil is applied directly to all surfaces of the food, it provides faster cooking (Singh, Varshney & Agarwal, 2016). In the frying method, the PAH content of the food differs according to the processing of the oil used. Repeated use of the used oil significantly increases the PAH content in the food. As a result, both the fried food and the air in the kitchen are contaminated with PAH with the gas released (Singh, Varshney & Agarwal, 2016).

7.4. Drying

Drying is one of the oldest methods applied in foods. The aim is to prolong the shelf life of the food by preventing bacterial growth in the food. This method can be applied to meat and meat products, vegetables, fruits, cereals and legumes. In this method, the air of the environment and the drying floor are among the factors that increase contamination during the drying phase (Singh, Varshney & Agarwal, 2016).

7.5. Steaming

Steam cooking method is the method of cooking food as a result of transferring heat from steam. Since pyrolysis is minimized in this method, PAH formation is also low. This method is the most important cooking method that supports low PAH formation. In this method, it was determined that PAH was formed 35 times less than the grill method, 24 times less than the smoking method, and 15 times less than the roasting method (Chen and Lin, 1997).

8. PAH – Impact on Health

The effects of PAHs on human health depend on the PAH concentration and exposure time of the contact contamination (Buha, 2011). The health of the individual is also related to other factors such as basic health status, genetic characteristics, environmental factors, age and gender before exposure (Buha, 2011). Epidemiological studies have shown that combinations of PAHs and duration of exposure affect basic health (ATSDR, 2011). Naphthalene, which is frequently found in potable water from PAHs that enter our lives in many ways, binds to molecules in kidney, liver and lung tissues with covalent bonds and increases the toxicity rate with this method (Sudip, Om & Rakesh, 2002). In addition, it is known to cause eye defects and hemolytic anemia (Srogi, 2007). In the short term, many symptoms such as vomiting, nausea, diarrhea, abdominal pain, headache, excessive sweating, and tachycardia can be seen with high exposure (IPCS, 2002). At the same time, mild irritation and dermatitis can be observed on the skin after high exposure in a short time (CDC, 2009). In addition to all these, urban air pollution and PAHs taken with food have negative effects on the immune system (Liu et al, 2013). In addition, both short-term and long-term effects of Acenaphthene caused adverse effects on the immune system and skin diseases, while PAHs such as Benzo (a) anthracene caused tumors as a result of animal exposure (Kim et al, 2013).

9. Conclusion and Recommendations

- ✓ In cases where the smoking method is not applied consciously, it is inevitable that undesirable compounds will occur in the food to be consumed. These hydrocarbons can accumulate in the human body over time and cause important health problems, especially breast, pancreatic and stomach cancer, where they can have mutagenic and carcinogenic effects.
- ✓ Instead of the traditional smoking techniques that are frequently used in our country; Smoked technology should be used in systems where smoke aroma, oven temperature and humidity inside the oven can be controlled, and the cooking smoke can be prepared outside the oven and then filtered and delivered to the nutrients in the modern oven. In this way, consumption of richly flavored incense products will not pose a health problem, and it will be possible to produce standard smoked foods similar to the practices in European countries.

- ✓ In addition, it should be preferred to bake and boil foods of animal origin (fatty meat) instead of grilling them by placing them on coal or direct fire. Pure charcoal should be used as fuel type. When grilling, the cooking process should be done on embers after the flames are extinguished, not on the burning fire.
- ✓ The distance between meat and fire should be at least 7 cm, ideally 10-15 cm.
- ✓ Measures should be taken to prevent air, soil and water pollution and inspections should be carried out effectively.

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Pandemics

Cihat Furkan Göv
Hülya Çelik¹

1. INTRODUCTION

The emergence and spread of infectious diseases with pandemic potential has occurred regularly throughout history. Many infectious diseases that lead to pandemics are caused by zoonotic pathogens transmitted to humans due to increased contacts with animals through breeding, hunting and global trade activities. Understanding the mechanisms of transmission of pathogens to humans allowed the creation of methods of preventing and controlling infections. For centuries, the implementation of public health measures such as isolation, quarantine and border control have helped control the spread of infectious diseases and maintain the community structure. Global friction programs of waterborne pathogens, vector-borne diseases and zoonotic spreads in the animal-human interface are of primary importance for rapidly detecting the emergence of infectious threats. In the event of a pandemic, new technologies are needed for rapid diagnostic tests, contact monitoring, reuse of drugs, biological markers of disease severity, and new platforms for the development and production of vaccines. In this research project, we review the major pandemics that have affected humanity throughout history, such as plague, cholera, influenza and coronavirus diseases, how they were contained in the past and how these diseases are managed today.

Virüs

The virus is a small agent that causes infection with only one of the deoxyribo nucleic acid or ribonucleic acids packaged in the protective sheath, which consists of a protein or complex structure. Viruses are not considered alive because they need host cells to grow and multiply. Since viruses do not independently have proliferation mechanisms and molecules, they depend on host cells for reproduction and are considered parasitic cells. Viruses often have a limited host range that they can infect, and viruses can only infect certain bacteria, plants or animals. Viruses depend on the metabolic molecules of the host cell. Viruses alter the host's normal metabolic reactions, allowing new virus particles to form. In order for viruses to survive, they must infect a cell and synthesize their own viral proteins and genetic material by capturing the mechanisms of the cell (Takz,2019).

Ways Viruses Are Transmitted

- Direct contact
- Injection of blood or contaminated liquids
- Tissue transplants
- Respiratory tract
- Fecal-oral pathway

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Factors Affecting The Ways Viruses Are Transmitted

- Source of the virus
- Resilience of the virus
- Environmental conditions
- How the virus reaches the target tissues of the body (Basustaoglu,2010).

Proliferation of Viruses

When viruses encounter a host cell that can multiply by entering the body of the living creature, they activate their proliferation mechanisms. When the virus encounters a suitable host cell where it can multiply, it immediately clings to it. After viruses cling to the host cell, certain enzymes are used to break down the cell membrane and transfer genetic material into the cell. This genetic material can be a chain of ribonucleic acid or deoxyribonucleic acid, which contains information that provides the synthesis of proteins necessary to initiate the process of developing new viruses. The genetic material of the virus captures the mechanisms of the cell through enzymes belonging to the virus. The mechanisms of the cell are used to create new genetic material belonging to the virus and to create protein sheaths. New viruses in the cell usually come out when the cell breaks down. When the cell disintegrates, the cell dies. This cycle, which viruses use to multiply and cause the cell to die, is called a lytic (disruptive) cycle.

Viruses with membranes on protein bristles do not choose the way to break down the body in order to multiply. Unlike other viruses, such viruses pass through the cell membrane and enter the cell. These viruses, which enter the cell as a whole, multiply and then come out without breaking down the cell. Once the virus enters the cell, it stores its genetic material as part of the cell deoxyribonucleic acid. In this way, the virus continues to transfer genetic material to the baby cells every time the cell multiplies without revealing its presence to the cell. It can secretly transfer the genetic material of the virus from cell to cell without causing any signs of disease, and suddenly the virus starts the cycle of proliferation. This cycle is called the lysogenic cycle (Freudenric and Kiger 2020).

Pandemi

The word pandemic is derived from the ancient Greek words pan and demos, and 'pan' means 'demos' and people mean (Pandemic). Wikipedia, free encyclopedia,2022).

The effect is not a specific country or region; Infectious diseases that are effective all over the world are called pandemics. According to the World Health Organization, the pandemic is a rapidly spreading and unseasonal infectious disease in which the human population does not have immunological resistance (Qiu et al.2017).

General Characteristics of Pandemics

- Appearance in various geographical areas
- The emergence of many diseases that can be a precursor to infectious diseases
- Caused by mutated viruses
- Inadequate or no social immunity
- Formation of diseases resulting in death
- Causing changes and measures in the lifestyles of society
- Spreading without seasonal conditions (Özkoçak et al. 2020).

Major Pandemics in Human History

Amvâs Tâ:

The plague, which occurred in the Palestinian settlement of Amvâs during the reign of Omar, spread rapidly and was also seen in the territory of Syria and Iraq. Syria was the most common settlement of the epidemic.

The plague of cucumbers spreading in Amvâs is the most common of the plague species. It is defined by the Arabs as a blessing.

Since it appeared in Amvâs, the epidemic was called Amvâs tuânu.

There are three types of this plague, bubonic, septicaemia and pneumonic, caused by a bacterium called *Yersinia pestis*, which passes from one rodent infected with fleas to another. Tâun is a bubonic species also known as cucumber, which is one of the plague species.

The bacterium *pestis*, which enters the human body with the bite of fleas, causes the lymph node to become inflamed and stretched.

In the later stage of the disease, with the growth of lymph nodes, inflamed and severe pain-causing bulges appear in the body

Swelling is more common in the armpits, behind the ear, earlobe and groin areas where the skin is thin in the body

Swellings can be red, yellow, black in color. In the mildest type of the disease, the silus is red in color. With the increase in disease severity, the swellings turn yellow and then black.

In the Middle Ages, medical examiners did not know much about the treatment of taun disease. If the swellings in the bodies of those infected with Taun's disease spontaneously exploded without intervention and the blood and pus in the boil emptied out, the patient had a chance of recovery. Otherwise, patients usually die within three days.

According to the imam shafi, he thought that the patient would be healthy by applying the ointment obtained by crushing the violet flower in the treatment of taun disease or by boiling the violet and containing it in the patient (Genghis,2020).

Black Plague

The disease was seen between 1347 and 1351. The disease is called black plague because of the appearance of black swelling in the body (Glitter, 2020).

Yersinia pesticide is the cause of the plague epidemic that has killed many people in Europe (Jarus,2020).

Doctors who treated the plague also wore some special costumes. The protective costume consisted of a polished fabric coat, a beak-shaped mouthpiece usually filled with herbs, straw and spices, and a mask with a glass eye part. These doctors studied ways to treat blood draw, frog or leech (Byfield, 2010).

Cholera

Cholera is an infection whose factor is *Vibrio Cholerae*, which causes the body to lose high rates of water, showing signs of diarrhoea that starts suddenly and suddenly intensifies. When the bacteria, which is resistant to stomach acid, is taken into the body at high rates with liquid, the bacteria escapes stomach acid and reaches the intestines, causing infection (Undored and Altinbas 2020).

The outbreak was seen in Japan in 1817, Moscow in 1826, Berlin in 1831, Paris and London, Hamburg in 1892 and Canada. Many people have lost their lives in these outbreaks. Treatment for cholera, which has a high mortality rate, is quite simple.

Cholera patients recover in a short time with oral fluid treatment. Salt and glucose mixtures are given to eliminate the lack of electrolytes such as sodium, potassium, chlorine and bicarbonate lost in the treatment and to provide the necessary energy in patients with nutritional problems. In patients whose condition is too severe to meet the nutritional needs, the mixture is applied into the vein. In even more serious and urgent cases, tetracycline and tetracycline-like antibiotics are used (Aslan,2020).

Spanish Flu

The influenza epidemic, a subtype of H1N1 virus, was the cause of the H1N1 virus between 1918 and 1921. The surprising thing about the epidemic is that it affects more healthy young adults than the weak, the elderly and children. The reason the outbreak is called spanish flu is because Spain did not take part in World War I and there was no mention of the outbreak in other European states due to censorship, and the Spanish press brought up the outbreak for the first time (Aslan,2020).

Doctors used vaccines against the hemophilus influenzae and pneumococcal species to treat the outbreak. They gave oxygen to the patients. Although they resorted to ancient applications such as bleeding the patient, they could not find a definitive treatment in every intervention.

In order to reduce the spread of the epidemic, measures such as quarantine, isolation, mask use, closure of public areas were taken but were unsuccessful.

The vaccines and drugs used in the treatment were not effective. Although blood transfer from recovering patients to new patients was a silver lining, there was no definitive solution (Temel, 1918).

Hong Kong Flu

The disease was first seen in Hong Kong. It is the influenza epidemic that caused the deaths of nearly 1 million people between 1968 and 1969.

The disease-causing type A H3N2 virus is derived from the H2N2 virus with an antigenic deviation called the merger of many subtypes of the virus to produce different viruses than the original.

The new type of virus carries the genes of H2N2, one of the bird flu virus variants (Aslan, 2020).

Symptoms of Hong Kong flu include fever, muscle pain, weakness. These symptoms usually lasted 4-6 days. The mortality rates of children and elderly patients with Hong Kong flu are high. Although a vaccine for the virus was produced, it benefited after the outbreak affected people on a global scale (Hsieh et al.2006).

Smallpox

Variola virus is the cause of smallpox. Variola virus is in the Poxviridae family. When a case of smallpox is encountered, it is obligatory to be reported internationally. Smallpox falls into the group of infectious diseases among the types of diseases (Özer and Suna 2020).

The first vaccine produced in history is the smallpox vaccine (Badur,2020).

In 1966, as a result of the campaign led by the World Health Organization, the use of smallpox vaccines was mandatory worldwide. The World Health Organization has reported that the infection that causes smallpox worldwide has ceased permanently. The World Health

Organization has recommended that the smallpox vaccine be removed from routine practice after the end of the disease. Smallpox is the only disease that has ever ended among human infectious diseases (Özer and Suna 2020).

Swine influenza

The cause of the disease is a virus in the family Orthomyxoviridae. The other name of the disease is "swine influenza virus", i.e. SIV.

Known swine flu types are usually Influenzavirus A, which is rarely type C influenza virus.

Swine flu is constantly changing its structure to spread among humans.

In 2009, an outbreak of swine flu was observed. The outbreak, which began in 2009, originated in Mexico and affected 191 countries worldwide. The outbreak that started in Mexico was caused by H1N1, a subspecies of influenza A virus.

The disease is transmitted to humans from those who come into contact with poultry and pigs closely and without protective clothing. When the meat of the sick animal is cooked under favorable conditions, the disease is not transmitted (Kara, 2010).

Symptoms of the disease are fever, cough without phlegm beats, unbearable pain throughout the body, fever, weakness, shortness of breath and sudden symptoms with a temperature of 38.30 °C. Sudden symptoms are high fever, pain, ache, diarrhea, which develop between 3-6 hours. Although these symptoms vary from person to person, they can usually vary between 4-7 days.

Oseltamivir or zanamivir drugs used to treat flu can be used in the treatment. These drugs should be used within the first 2 days after the onset of symptoms. Drugs should be administered under the supervision of a doctor and used in accordance with risk groups. Paracetamol should be used instead of aspirin as a painkiller. It is not recommended to consume medicines when going abroad for pandemic prevention purposes (Sanli,2010).

Bird Flu

Bird flu is a deadly disease transmitted from animals with the virus. Other names of the disease are Avian Influenza, Pestis Avium, Bird Flu, Avian Flu, Chicken Plague. The type of virus that is deadly to humans is H5N1 (Aslan,2020).

Bird flu is an infectious disease caused by influenza viruses found in birds. Disease-causing viruses are carried in the intestines of birds, are found in their feces and often do not show symptoms of the disease.

The virus that causes bird flu has been proven to be transmitted between pigs and humans, between chickens and humans, but there is no certainty about whether there is transmission between wild birds and humans (Goldrick and Goetz 2007).

Symptoms of the disease include fever, cough, sore throat, muscle aches and pain in the lung. Cases with pneumonia and respiratory distress are at risk of death. The disease is called blue death because the body begins to turn blue when not enough oxygen enters the cells.

There is no definitive treatment for the disease. Protection is provided by vaccination. Animals with disease factors need to be killed. Due to antigen changes in influenza virus, the activity of the virus should be monitored and the content of influenza vaccines should be updated every year (Aslan,2020).

Ongoing Pandemics

. Hiv/Aids

The hiv virus is the cause of the disease. Hiv virus is known as the human immunodeficiency virus. Hiv virus affects the mechanisms of the immune system in humans, causing the system to collapse. The disease that hiv virus is a factor in is called AIDS. AIDS is known as acquired immunodeficiency syndrome. AIDS is an infectious disease in the zoonotic group (Undored and Altinbas 2020).

HIV virus, which is a factor in AIDS disease, is the virus that causes the collapse of the immune system by affecting cells involved in vital functions such as T cells, macrophages and dendritic cells, which are auxiliary cells in the human immune system (Özer and Suna 2020).

Ways hiv/aids are transmitted

HIV virus, which is a factor in AIDS disease, is found in the blood of people infected with the highest rates. Genital secretions, phlegm, breast milk, saliva, tears and cerebral spinal fluid may also be a disease factor. The most important factors that are effective in the transmission of the disease factor are blood, genital secretions and breast milk.

Stages and Cemptions of HIV Infection

Primary Infection

Hiv virus, which is a factor in AIDS disease, first acts on macrophage/dendritic cells in submucosal layers;

Approximately 87% of people infected with HIV 2-4 weeks after entering the body show flu-like symptoms. . People infected with this stage have high rates of HIV in their genital fluids and have a high amount of transmission (Lewthwaite and Wilkins 2009).

Serokonversiyon

This period, known as seroconversion, is the stage at which the body begins to produce antibodies to the virus (Card et al.2008).

Asymptomatic Infection

The stage of asymptomatic infection is known as the latent period. At this stage, the disease progresses quite insidiously. It does not show symptoms or symptoms. The duration of this period varies from person to person. The reason for the difference in duration is due to the difference in the health status and immune system functions of the person (Lever,2009).

Advanced HIV Infection/ Clinical AIDS

It is the stage where there is a significant reduction in immune system mechanisms, increased proliferation of HIV virus, cessation of antigen production and increased susceptibility to infections (Hazenberget al.2003).

HIV /AIDS Treatment

The aim of the treatment, known as antiretroviral, is to suppress HIV replication and increase immune functions by using drugs that control the proliferation of HIV virus, to keep the life expectancy of infected people at normal level and to normalize the quality of life. Effective viral suppression has a positive effect on the general health of the community by reducing the risk of disease transmission among individuals (Akgül et al. 2018).

Drugs Used to Treat HIV/ AIDS

Nükleozid Revers Transkriptaz İnhibitörleri (NRTİ)

Nucleoside Revers Transcriptase Inhibitors interact with the catalytic region of the revers transcriptase enzyme found in the HIV virus, a cause of AIDS disease (De Clercq,2007).

Nucleoside Revers Transcriptase Inhibitors have analogues 2',3'-dideoxynucleosine (ddN), and the effects of drugs in this group are similar. After nucleoside Revers Transcriptase Inhibitors enter the cell, they turn into 5'-monophosphate (ddNMP), 5'-diphosphate (ddNDP) and 5'-triphosphates (ddNTP), respectively. The transformation process occurs before ddNMP acts as a competitive inhibitor/alternative substrate of normal deoxynucleoside triphosphate (dNTP).

As a compelling inhibitor of the normal substrate, ddNTP inhibits the merging of the normal substrate with the DNA chain and participates in this chain as an alternative substrate in the form of ddNMP. Thus, it acts as a de-chain terminator (De Clercq,2009).

Non-Nükleozid Revers Transkriptaz Enzim İnhibitörleri (NNRTİ)

The first two classes of Non-Nucleoside Reversal Transcriptase Enzyme Inhibitors are known as 1-[(2hydroxytoxy)methyl]-6-(phenylthio)timin-structured compounds i.e. 4,5,000 with HEPT6.7-tetrahydro-5-methylimidazo[4,5,1-jk][1,4]benzodiazepin-2(1H)-on, i.e. TIBO derivatives (Pauwels et al. 1990).

Compounds of this nature primarily interact with the allosteric region of HIV-1 revers transcriptase. The drugs in this group are also called specific inhibitors of HIV-1 (De Clercq,2004).

Protease Inhibitors (PI)

Protease Inhibitors carry a hydroxyethylene skeleton opened by the HIV protease enzyme, which resembles a peptide bond but cannot be opened itself. The mechanism of action of drugs in this group prevents HIV protease from performing its normal function; thus, the proteolytic process of precursor viral proteins to mature viral proteins is prevented (Madruga et al.2007).

Fusion Inhibitors (FI)

Hiv virus, which is a factor of AIDS disease, creates fusion with normal cells thanks to gp41 in the membrane, transferring its viral RNA and viral enzymes to normal cells. Enfuvirtide, a physio inhibitory, was produced to prevent this fusion (Matthews et al. 2004).

The mechanism of action of drugs in this group is to prevent the virus molecule from fusing with the outer cell membrane (De Clercq,2009).

Co-receptor Inhibitors (CRI)

Co-receptor inhibitors must enter the target cells to show their effects, and the correlators used in M (macrophage)-tropical and T (lymphocyte)-tropical HIV lineages) should interact with CCR5 or CXCR4. The mechanism of action of drugs in this group is the interaction of the co-receptor and viral glycoprotein gp120, the interaction of gp120 with the CD4 receptor and the fusion of gp41 and the outer cell membrane (Westby and van der Ryst 2005)

The only CRI drug approved for use is maravirocytose (Perros, 2007), an antagonist ccr5.

Integrators (II)

The enzyme integral has attracted attention in the development of drugs against hiv virus. The first clinically approved integral inhibitor (II) is raltegravir. The effect of raltegravir is on thread transfer. This is the effect of elvitegravir. Elvitegravir is effective in reducing viral loading in people infected with HIV (Sulukan et al. 2009).

Covid-19

Coronaviruses are single-chain, positive polarity, enveloped RNA viruses in the Coronavirinae subfamily of the Coronaviridae family in the Nidovirales class. The reason these viruses are called coronaviruses is due to the latin word "corona", which means "crown" due to the bar-shaped extensions found on the surface of the virus (Chen et al.2020).

The four structural proteins found in coronaviruses - the S (Spike) protein, the E (Envelope) protein, the M (Membrane) protein and the N (Nucleocapsid) protein -- are responsible for the formation of mature virus particles (virions) and the emergence of infection. The S protein is responsible for the formation of spikes on the surface of the coronavirus. The protrusions on the surface are responsible for the virus's adhesion to the host cell receptors and its entry into the cell. The M protein, which allows the virus particle to form with three different transmembrane regions, to allow the virus to curve the cell membrane and bind to the nucleocapsid. Protein E is involved in the packaging and release of the virus and viral pathogenesis. The N protein contains regions connected to the viral genome in the RNA structure (Alpine and Unal 2020).

Ways Covid-19 Is Transmitted

The main way of transmission of coronavirus is the droplet path. The virus found in respiratory secretions can make the healthy individual sick by direct contact with the mucous membrane, while the infected individual coughs, sneezes or speaks through droplets. In finding a new host for the virus, the contact of another individual is effective with the droplets that the sick individual emits through coughing and sneezing (Gürsu,2020).

Stages and Symptoms of Covid-19 Infection

The severity of infection is divided into 5 groups: asymptomatic infection, mild form, moderate form, severe infection, critical infection.

Asymptomatic Infection

It is defined for cases that have tested positive for coronavirus and do not show clinical symptoms. Asymptomatic infection plays an important role in social transmission. This is due to the absence of clinical symptoms in those infected. Babies and children who do not show clinical symptoms play an important role in interpersonal transmission.

Mild Form Symptoms

Fever, fatigue, muscle pain, cough, sore throat, runny nose, sneezing, symptoms of acute upper respiratory tract

Symptoms of Medium Form

Pneumonia, usually fever and cough, wheezing in some cases

Symptoms of Severe Infection

Dyspnea, central cyanosis, decreased oxygen saturation, hypoxemia

Critical Signs of Infection

Acute respiratory distress syndrome, respiratory failure, multiple organ failure (Happy, 2020).

Covid-19 Treatment

The main goal in the treatment of coronavirus is supportive and symptomatic treatment. First of all, the infected persons should be removed from the community and quarantined. Cases with mild symptoms should be managed by home counseling without going to health centers. In

treatment, the body's water and electrolyte level should be kept to a certain extent, nutrition should be taken care of, fever and cough should be checked. In cases where the body does not receive enough oxygen, the face mask, high-flow nasal cannula, oxygen should be provided through non-invasive ventilation. Mechanical ventilation and oxygen support may be required, and kidney replacement therapy may be required in some patients. Antibiotics and antifungal should be used in cases with two or more infections in the body at the same time (Jin et al. 2020).

Vaccines are the most effective way to prevent epidemics in the form of infections. This is because it takes both costly and a long time to develop drugs suitable for treatment. Vaccines are both a way to be used in the prevention of global epidemic diseases (DG et al. 2020).

Drugs Used to Treat Covid-19

Klorokin (CQ), Hidroksiklorokin (HCQ)

Although chloroquine has not been approved by the FDA in the treatment of coronavirus, it has allowed use if doctors deem it appropriate. The FDA has not approved any therapeutic agents or drugs for the treatment and prevention of coronavirus. Clinical efficacy and safety of chloroquine in the treatment of coronavirus is unclear. Since doctors are successful in the treatment of different diseases, they have approved the use of chloroquine.

Hydroxychloroquine is a chloroquine-derived drug. The mechanism of action of the drug inhibits the enzyme polymerase, preventing the merger by interfering with the glucocytion of the virus and cell receptors by increasing the ph of the place where the intracellular digestion of the substance formed and taken by the combination of phagosome and lysosome occurs.

Hydroxychloroquine is recommended in patients who will be monitored with outpatient treatment and treated by hospital in the coronavirus treatment guide of the Ministry of Health in our country.

Favipiravir

Favipiravir RNA viruses have an effect by inhibiting the enzyme RNA-dependent RNA polymerase. Favipiravir is an antiviral drug. Favipiravir forms favipiravir ribofuranosyl-5B-triphosphate, an active form of the body. For this activation process, favipiravir intraselecular phosphoribosylation occurs. Favipiravir ribofuranosyl-5B-triphosphate is a substrate of RNA-dependent RNA polymerase and inhibits RNA polymerase enzyme activity. Since the RNA-dependent RNA polymerase enzyme is found in many RNA viruses, favipiravir expands its spectrum by affecting this enzyme.

Remdesivir

Remdesivir is in the antiviral drug group. It is used in the treatment of Ebola and marburg virus infections. This drug is the precursor to nucleotide analoma, which is metabolized into the intracellular adenosine triphosphate analogue, which inhibits the enzyme viral RNA polymerase. Remdesivir is a wide-spectrous drug. It has a preventive and therapeutic effect in non-clinical studies against coronaviruses.

Lopinavir/ritonavir(LPV/R)

Lopinavir, a combination of ritonavir, is used to treat HIV infection. It is a protease inhibitor, one of the drugs used to treat AIDS. The enzyme protease plays a role in the processing of coronavirus polyprotein. Lopinavir and rheaavirin have contrasting effects against coronavirus separately and together.

Nitazoksanid

The drug nitazoksanid has strong in vitro activity against coronavirus. Apart from the coronavirus, this drug has broad-spectrum antiviral activity against influenza, respiratory syncytial virus, parainfluenza, rotavirus and norovirus. The mechanism of action of the drug is associated with the pathways regulated with the host during the proliferation of the virus.

Tosilizumab

The effect of the drug tosilizumab is caused by monoclonal antibodies due to membrane or inhibiting soluble interleukin-6 receptors (rIL-6). This drug has been approved for use in the treatment of rheumatoid arthritis patients.

İvermektin

Ivermectin is a broad-spectrum FDA-approved drug effective against parasites. It has had an effect on many viruses in studies on the drug. The mechanism of action of the drug ivermectin inhibits the nuclear intake of the host and viral proteins. Ivermectin both acts on some RNA virus infections and has an effect on some DNA viruses. In studies on the drug, the coronavirus has a reducing effect on its effectiveness (Mutlu et al. 2020).

Factors Affecting Viral Pandemics

Vaccines

Viral pandemics can be effectively prevented with inexpensive and effective vaccination strategies. Vaccines protect against pathogens by stimulating primary defense mechanisms.

Vaccination is the process of applying a small amount of an infectious agent, such as viruses or bacteria, to stimulate the immune system to recognize and respond to this agent. Vaccines are often used to prevent infectious diseases. Vaccines are administered by injection. Vaccines are often designed to stimulate the production of antibodies, proteins that neutralize foreign invaders in the body.

To improve the specific immunological system through vaccination, the human body can produce white blood cells (B and T cells) that can fight the pathogen (especially) in response to the immune system, allowing the adaptive immune system to grow.

Since the disease is mostly a new type of virus in virus-induced pandemics, it is impossible to produce a vaccine before the pandemic begins.

In the production of vaccines against pandemics, the disease factor is primarily stated. After the disease factor is determined, it is replicated in the cultural environment and the vaccine is produced. The spread of pandemics increases in the time spent in vaccine production. The vaccine produced is specific to the disease factor determined. Vaccines are effective on the black virus. The treatment of the infected person with the vaccine is limited (Şahin and Demir 2020).

Antiviral Drugs

Antiviral drugs are used in virus-induced infections. It is used in the prevention and treatment of virus-induced infections. In viral pandemics, the treatment of the disease is either through the application of vaccines or the use of antiviral drugs to prevent the disease before the disease occurs (Carrasco et al. 2011).

Virus-Related Protein Use

The protein associated with the virus contains anti-idiotypal antibodies, natural ligands of the receptor and anti-receptor antibodies. The mechanism of action prevents the virus from binding to host cell receptors and entering the cell before the viral infection begins.

Reverse Transcription Inhibitors/ Revers Transcription Inhibitors

Nucleotide or nucleoside analogues, similar to the building blocks of ribonucleic acid or Deoxyribo nucleic acid, are used as antiviral drugs. Since the mechanism of action is not 3'OH in analogues, viral Ribonucleic acid or Deoxyribo blocks enzymes that synthesize nucleic acid.

Antisense

When the virus genome becomes active in the host cell, it produces mRNA molecules that play a role in the synthesis of viral proteins. mRNA production begins by proteins. Antisense drugs are made from short segments of DNA or RNA that can bind to their viral genomes as a complement and prevent transcription of the viral genome.

Ribozim

When the virus genome becomes active in the host cell, it produces mRNA molecules that play a role in the synthesis of viral proteins. mRNA production begins by proteins. Antisense drugs are made from short segments of DNA or RNA that can bind to their viral genomes as a complement and prevent transcription of the viral genome.

Protease Inhibitors

Protease inhibitors developed for the treatment of HIV/ AIDS and hepatitis C are widely used in the treatment of these diseases. Some viruses break down viral protein chains after translation with the enzyme protease within the target cell, creating virions and spreading them in cells that are not infected with disease factors. Protease inhibitors take effect by inhibiting the enzyme viral protease.

İnterferonlar

Interferons suppress the disease-causing virus factor by strengthening the body's immune system instead of attacking the disease-causing virus factor. Interferon viruses prevent their proliferation. Interferons are signaling proteins produced due to the presence of several viruses in the body. Interferons are cytokines synthesized by cells in viral infection. It shows its effect by enabling communication between cells to strengthen the immune system against disease-causing factors. Interferons have opposite effects on the proliferation of viruses by protecting cells from virus infections.

Conventional Plasma Treatment

Conventional plasma therapy, which is used to treat and prevent epidemic diseases, is the treatment of the disease by taking blood from people who have suffered from the disease, separating antibodies and giving them to sick people (Şahin and Demir 2020).

3. RESULTS AND RECOMMENDATIONS

Pandemic epidemics that are effective worldwide. An epidemic pandemic not only is it widespread and the mortality rate is high, but it is also contagious must be high. Affecting human history pandemic. Although various methods have been applied in the treatment of related diseases for centuries, there is no definitive treatment method. For pandemic there is a unique way of treating the disease. These symptomatic and supportive treatment, lac treatment and vaccination of healthy people pandemic to reduce the effects. Pandemic investments in preventing factors increase, Ar-Thanks to the development of ge studies, we may face in the future pandemic we strengthen our hand in their dangers.

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Hemorrhoidal Disease & Treatment

Hüseyin Kerem TOLAN

Introduction

Hemorrhoids are anatomical structures of the anal canal which contribute to the continence mechanism. Due to various reasons they may become enlarged, engorged and thus symptomatic. The often encountered symptoms are bleeding, mucosal soiling and pruritus mostly during and after defecation but they may also occur spontaneously. A colorectal surgeon must comprehend this clinical entity well since it comprises a significant amount of outpatient clinic admissions and other benign proctological conditions can easily be mistaken as hemorrhoids by both patients and health professionals.

Anatomy

The upper anatomical part of the surgical anal canal is called Anal Transitional Zone (ATZ) where cushion like venous plexuses reside in its subepithelial space, hence the name Corpus Cavernosum Recti (CCR) (Fig. 1). This plexus is not continuous and often three distinct cushions (left lateral, right anterior and right posterior in supine position) are prominent, which are called hemorrhoids. These veins tend to be tortuous in nature and with their surrounding smooth muscle fibers and connective tissues, they are also referred as anal glomerula (1). They are anatomical structures. The integrity of these plexuses are maintained with the muscle fibers originating from the Internal Anal Sphincter (IAS) and Conjoined Longitudinal Muscle (CLM). Superior Rectal Artery (SRA) is the main arterial source while the venous drainage is maintained by hemorrhoidal veins.

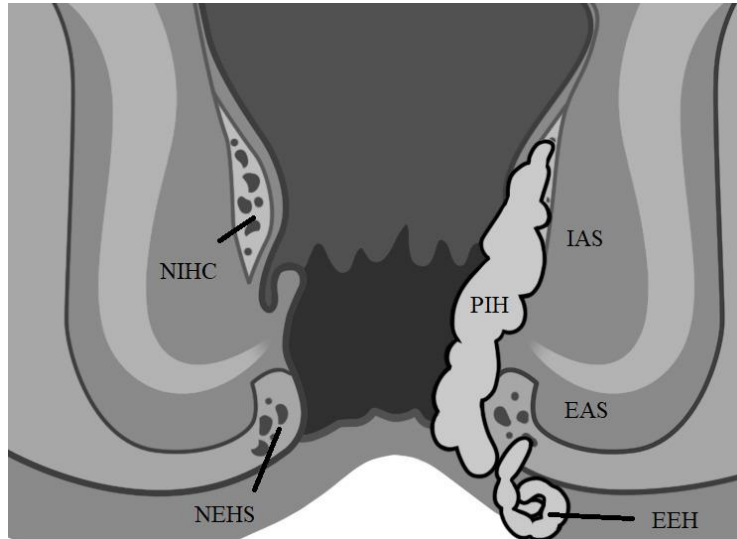


FIGURE 2. The Illustration of Sphincters and Hemorrhoidal Structures. IAS:Internal Anal Sphincter, EAS:External Anal Sphincter, PIH:Prolapsed Internal Hemorrhoid, EEH: Engorged External Hemorrhoid, NIHC:Normal Internal Hemorrhoidal Cushion, NEHS: Normal External Hemorrhoidal Structure

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Epidemiology and Etiology

Although it is challenging to assess the true prevalence of the hemorrhoidal disease due to the fact that most of the individuals remain asymptomatic and therefore do not seek medical treatment, the contemporary literature reports a disease burden of 10% among the general population with a slight female predisposition (2).

Various factors contribute to the pathogenesis. For this reason it is hard to define a single mechanism to be the underlying cause. However, almost every factor contributing to the development of this disease ultimately result in either impaired venous drainage or destruction of the connective tissue within the cushions (3). In most cases these two main alterations overlap. Studies demonstrated an increased metalloproteinase activity, namely MMP-9, and neovascularization due to overexpression of VEGF and TGF- β in clinically significant hemorrhoidal cushions (4,5). Additionally, the resting anal canal pressure is found to be elevated compared to healthy individuals in patients with hemorrhoidal disease (6). Ultimately two distinct theories are generally accepted;

1. Hyperplasia Theory: Impaired venous drainage of CCR due to the increased IAS pressure which results in prolapsus of the cushions into the anal canal.

2. Anal Sliding Theory: Submucosal smooth muscle destruction as a result of increased intraabdominal pressure, which again leads to prolapsus. Chronic constipation, chronic cough, pregnancy and numerous other entities can result in increased intraabdominal pressure.

Interestingly, no association between increased hemorrhoidal disease prevalence and liver cirrhosis could be shown, even though cirrhosis clearly increases the intraabdominal pressure (7). Nevertheless, rectal varices develop in most of the cirrhotic patients and can be mistaken as hemorrhoids.

Classification

Hemorrhoids which originate from the ATZ are described as internal hemorrhoids. Occasionally, hemorrhoids may also occur below the dentate line and are referred as external hemorrhoids. Contemporarily, Goligher Classification is still applied to grade internal hemorrhoids (8). According to this classification four distinct grades can be distinguished (Fig. 2);

Grade I: Slightly enlarged cushions with no visible prolapsus.

Grade II: Cushions prolapse and become visible during defecation or straining but reduce spontaneously.

Grade III: Prolapsed cushions which can be reduced manually.

Grade IV: Irreducible cushions.

Despite being routinely applied in clinical setting worldwide, this classification lacks some important aspects of this disease, such as circumferential hemorrhoids, acute thrombosed hemorrhoids, mucosal prolapsus, etc. Some new classifications are proposed to address this issue. Unfortunately, the data in this regard are sparse and thus no optimal classification can be applied yet.

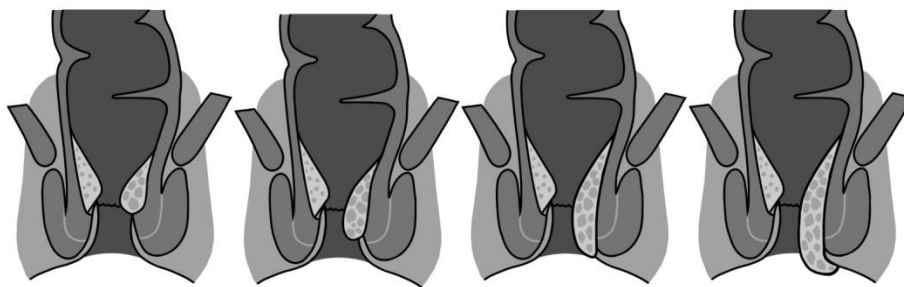


FIGURE 2. Goligher Classification of Hemorrhoids (Grade increases from left to right)

Clinical Presentation and Diagnosis

A hemorrhoidal disease presents itself most commonly with rectal bleeding, prolapsus, itching and pain. Pain is more common in external hemorrhoids. However, when the prolapsed hemorrhoid also includes an external part or it gets thrombosed, pain may still occur. Itching is also very common, but if it is the only symptom the patient describes, the practitioner should focus on pruritus ani rather than hemorrhoidal disease. When pain is the sole symptom with no visible external hemorrhoids, anal fissure should be ruled out, because patients tend to misinterpret skin tags for hemorrhoidal prolapse. When a thrombosis occur within a hemorrhoid cushion, the pain onset is sudden and its course is relentless. Patients with a history of ongoing or intermittent diarrhea should be evaluated for inflammatory bowel diseases. One needs to keep in mind, that if the pain is not accompanied by a visible hemorrhoidal cushion, perianal abscess might be the underlying condition. Other anorectal pathologies such as fistulas, malignancies should also be ruled out.

Nowadays, it is recommended by the european authorities to evaluate the first 15 cm of the rectum with a rectoscopy, in case of a suspicion of hemorrhoidal disease or even if the diagnosis is certain (9). A patient should undergo a complete colonoscopy if any predisposing factors to colorectal malignancies are present, such as increased age, family history, etc.

The physical examination can be performed in both prone-jackknife or left lateral position. A digital rectal examination is a must under any circumstances, except for the acute anal fissures, and the squeeze and resting anal canal pressures must also be evaluated during this examination.

Treatment

The initiation of a treatment only depends on whether the disease is symptomatic or not, regardless of its grade. In general, conservative treatment precedes and is favored over surgical approaches, except for the emergent cases, such as gangrenous hemorrhoids.

Conservative Treatment

First of all, the diet of a patient should be queried before commencing the patient on any medication. If the patients does not consume enough dietary fibre, it becomes very difficult to maintain a healthy bowel homeostasis. Therefore, it is recommended to consume 25 g daily fibre for women and 38 g daily fibre for men (10). The daily water intake should also be queried and if it is deemed insufficient, an adequate water consumption must also be ensured. With these dietary changes, constipation, the most common problem among patients with hemorrhoidal disease, can be overcome. Patients with chronic constipation may further continue suffering even under a strict fibre diet. In these patients, oral laxatives can be administered for a limited period of time (usually less than a week) (11).

Patients must also be educated about toilet habits. Prolonged defecation time and excessive straining are both associated with increased hemorrhoidal protrusion rates. The conditions related with the Obstructive Defecation Syndrome are beyond the scope of this chapter, however patients

can at least be advised f.e. not to bring a mobile phone to toilet, since this kind of habit clearly and inevitably prolongs the toilet stay. Patients should also be encouraged to have sitz baths 3 times a day for 10-15 mins with warm water. Applying warm water relaxes the IAS and thus facilitates venous return which alleviates the symptoms. A detachable shower head can also be utilized as a proxy for patients who can not perform sitz baths for various reasons.

Both flavonoids (phytonutrients which increase the venous tonus) and calcium dobesilate (a vasoprotective agent) are effective oral therapy options (12, 13). The recommended dose of calcium dobesilate for the acute setting is 500 mg twice a day, and the maintenance dose is 500 mg once a day. Any topical therapy, which results in decreased IAS tonus, alleviates pain or facilitates vasoconstriction, can theoretically be beneficial in treatment. Although no significant harmful effects have been reported in their utilization in hemorrhoidal disease treatment, the data regarding their efficacy in this setting are sparse. We prefer ointments over suppositories, since the latter is harder to maintain in the optimal anatomical position.

Surgical Treatment

Surgical treatments can be put into two main categories; Resection and non-resection techniques. Some of the non-resection techniques can also be performed in outpatient settings and are referred as office-based treatments. They include rubber band ligation, sclerotherapy, infrared coagulation and occasionally laser hemorrhoidoplasty. All of the office-based procedures target internal hemorrhoids, since the anatomical area is significantly less associated with pain. Non-resection techniques mainly aims to facilitate fibrosis around the CCR and thus fixation of the ATZ.

Sclerotherapy

This technique comprises of the injection of a sclerosing agent into the apex of a hemorrhoidal cushion. Various agents can be utilized, however ethoxysclerol or aliminium potassium sulfate and tannic acid (ALTA) are favored over the others (14). It is showed that ALTA is associated with less bleeding than phenol in almond oil (15).

The patient is positioned either in supine or prone-jackknife position and at the apex of each pedicle 1-3 mL sclerosing agent is injected depending on the agent and its concentration. Care should be taken to inject the substance directly to submucosa and avoid IAS, mucosa and CCR. This technique is relatively easy and painless. However, it is associated with high recurrence rates in the long term and repeat attempts are not encouraged due to risk of stricture. This therapy is effective in Grade I-II hemorrhoids.

Infrared Coagulation

This technique provides the same fibrosis and fixation mechanism with infrared energy instead of a sclerosing agent. Patients with low grade hemorrhoids are candidates for his procedure. A special, portable device is utilized. With the tip of the device, the apex is targeted and 3-4 applications can be done for each hemorrhoid. A recent prospective study favored this technique even over resection procedures (16). However, the data are sparse and the risk of local necrosis limited its use.

Rubber Band Ligation

The principle is that the apex of a hemorrhoid and the underlying rectal mucosa are strangulated with a band. Fibrosis and fixation occur consequently. The hemorrhoid is visualized with a proctoscope. Then the targeted cushion is pulled inside the device either with suction or a grasper and the band is deployed.

Multiple cushions can be addressed at the same time, although it is still advised not to apply bands to more than three cushions in one session (17). The procedure can be repeated at monthly

intervals. Rubber band ligation is effective in grade I-II hemorrhoids. Its role in grade III hemorrhoids is still an ongoing debate.

Band slippage and bleeding are common complications. Interestingly, bleeding can also occur 5-7 days after the procedure. Care should be taken, as in all surgical procedures for hemorrhoidal disease, not to overlook the signs of pelvic sepsis, which manifests itself at the early stage with urinary retention, pelvic pain and fever. However, pain is usually attributed to bands which are deployed too close to dentate line. In case of prolonged pain or in the presence of sepsis signs, either an anal examination under general anesthesia or computed tomography (CT) must be performed.

Laser Hemorrhoidoplasty

By using a 980-nm laser diode, the apex of the hemorrhoid is targeted, photo-energy delivered, arterial inflow diminished and consequently, fibrosis and shrinking are achieved (Fig. 3). The intraoperative positioning is the same as the others. The pedicle is grasped gently. Then the probe is inserted inside the pedicle above the dentate line and moved upwards. Three laser shots are made in the apex, two in the middle and the last one in the entrance area. Care should be taken not to damage neither IAS nor mucosa. After targeting each pedicle, we apply cold (mostly an icecube in a glove) for a minute to avoid hematoma.

This method is most effective in grade II hemorrhoids. The advantages are its short operation duration, less pain and low recurrence rates (18). However, despite being evaluated in a few studies, the data are sparse about its usage in grade III hemorrhoids. There is also a lack of information as to whether this technique is superior to other techniques. Therefore, the exact role of this method will be determined by future prospective studies.



FIGURE 3. Neo v Laser 1470 Nm, Diode Laser Machine. G.N.S neoLaser Ltd.©

Doppler-Guided Hemorrhoidal Artery Ligation

A special scope with a doppler device on its tip is inserted into the anal canal (Fig. 4). At the apex of the hemorrhoidal pedicle, its artery is identified and suture ligated. The ceasing of the arterial flow sound confirms a successful intervention. Once the arterial inflow is stopped, the rest of the mechanism is the same as with the laser method. Many surgeons also add a mucopexy by oversewing the pedicle caudally with the same suture till the dentate line and then knotting the first and the last thread together. The result is lifting of the pedicle, hence the name anal lifting.

This technique is associated with low recurrence rates, short operation times, especially in grade II-III hemorrhoids and when a mucopexy is done as well (19). Its main disadvantage can be pain. The rest of the complications are similar to the other above-mentioned procedures. However, data are sparse about this technique, too. Although there are some qualified trials about this procedure, more prospective trials are required to understand the true value of this method (20).



FIGURE 4. Comepa Angiodin-Procto ADL. Comepa Industries ©

Stapled Hemorrhoidopexy

This is a resection based technique performed by using a type of circular stapler. First the mucosa and submucosa of a circumferential line 4 cm above the dentate line is sutured continuously. Then the anvil of the device is introduced to the proximal of this line. The stapler is closed, with the hemorrhoidal pedicle is partially entrapped within, and fired. The excision of a mucosal doughnut allows lifting of the remaining hemorrhoidal pedicles, thus reduce prolapsus. Care should be taken not to involve rectovaginal septum, as this results in rectovaginal fistulas. The staple line should always be checked after firing the stapler, since bleeding occurs from this line and bleeding is the most common complication of this procedure.

This technique was first developed in 90's to present an alternative to conventional hemorrhoidectomy in grade II-III hemorrhoidal diseases. The literature suggests that it should not be undertaken in hemorrhoids which can not be reduced manually or which are thrombosed (21). Although this technique is associated with less pain compared with conventional methods, the recurrence rate is higher and one of the main downsides is that it reduces rectal compliance. Therefore many surgeons avoid using this technique more than once for the same patient. In our surgical practice, we tend to prefer this method less compared with the past, since our experience with other less invasive methods are more satisfying.

Excisional Hemorrhoidectomy

Excisional methods are basically reserved for Grade IV hemorrhoids or emergent situations, such as gangrenous hemorrhoids. They are still the gold standart in hemorrhoidal disease treatment. Recurrence rates are very low. However, they are associated with significant postoperative pain and stricture risk (22).

Except for Whitehead procedure, which is reserved for circumferential hemorrhoids, there are mainly three conventional excisional surgical interventions which are almost identical except for the wound closure at the end of the operation.

Basically, after positioning the patient, a combination of a local anesthetic and epinephrine 1:200,000 is injected in the perianal skin and inside of hemorrhoids. This allows less bleeding during

the procedure. The pedicle is grasped. The perianal skin is incised and the hemorrhoidal structure is dissected and lifted off the underlying sphincter structures until the apex, where the vascular pedicle is reached. We prefer using an electrocautery till this stage, however other options like bipolar devices or simply scissors are also viable. Once the apex is reached the pedicle is either stitch sutured or sealed with a bipolar device. Care should be taken to leave adequate skin-tissue bridges as the length of perianal skin incision circumference should not exceed the half of the total perianal skin circumference. Otherwise a postoperative anal stricture is likely to develop.

The above-mentioned steps are common. Once the excision is completed, Milligan-Morgan technique leaves the wound open to secondary healing. Parks technique recommends the closure of wound including the anoderm and Ferguson closes the perianal skin as well. Leaving the wound open exposes the patient to a slightly more stricture risk while suturing the wound may cause slightly more infection. However, these are not great differences and can be omitted. None of these methods are significantly superior to another and we believe it is up to surgeon's preference to tailor them for each patient individually.

Patients should be advised to continue using sitz baths, have daily shower with warm water and keep the area clean with water after defecation. Neither antibiotics nor anal tamponade are not mandated.

Whitehead excision mandates a circumferential incision along the dentate line. All hemorrhoidal structures are excised as well as excessive rectal mucosa. Then the remaining rectal mucosa is sutured to anoderm. This operation is highly effective as it removes all of the pedicles, but it is also associated with serious complications such as strictures, whitehead deformity, gas incontinence, etc (23).

Complications and Their Management

Anal Stenosis

The cause is the excision of too much anoderm. The clinical picture may turn out to be a mild one, which only requires laxatives. However, some patients may undergo dilation or even stricturoplasty (24).

Urinary Retention

This is a common complication of hemorrhoidal interventions, which are usually attributed to perioperative excessive IV fluids, inadequate pain relief and use of anal tamponades (25). The risk also increases if the patient has a urinary retention history such as benign prostate enlargement. To avoid this complication, the above-mentioned risk factors should be eliminated. Some surgeons also prefer using a urinary catheter, but this approach is still an individual practice rather than a scientific recommendation.

Pain

Anoderm is the sensible part of the anal canal. Therefore if an area below the dentate line gets operated on, pain occurs. However, despite some techniques being less related with pain, all surgical procedures can cause postoperative pain. Therefore, a routine commencement on postoperative analgesia is recommended. If the pain persists more than a week despite adequate analgesics or pelvic sepsis signs occur, an examination under general anesthesia must be performed.

Postoperative Hemorrhage

Postoperative bleeding occurs either in the immediate period (within 48 hours) or within postoperative days 7-10. Early bleedings are associated mostly with a technical error, such as overlooked stapler line bleeding. The cause of delayed hemorrhage is not certain, but it can be attributed to necrosis or sepsis of the pedicle apex. These hemorrhages rarely requires early surgical

intervention. Mostly an anal tamponade and close monitoring of the patient are sufficient measures. If the bleeding persists or causes hemodynamic instability, readmission to the operation theatre becomes a necessity.

Infection

This is a rare complication of hemorrhoidectomies. If it is suspected, antibiotics should be administered. If an abscess develops, it must be drained. A physician should always take the possibility of a pelvic sepsis and fornier gangrene into account, when evaluating a patient in regard of postoperative infection.

Fecal Incontinence

Hemorrhoidal cushions contribute mostly to gas continence. That means when they are excessively removed, a mild incontinence may occur. But this is usually a rare condition. Another contributor to fecal incontinence is damage to the IAS during surgery. A keypoint in addressing this issue is to evaluate the patient preoperatively in this regard. Especially parous women of advanced age are under risk. One should also keep in mind that stapled hemorrhoidopexy reduces the rectal capacity and thus may cause urge incontinence.

Special Clinical Pictures

Thrombosed External Hemorrhoid

The patient usually describes pain with a sudden onset. It is continuous and in some cases very severe. If the patient is compliant and the pain is manageable, a usual conservative therapy as described above should be commenced. If the pain is severe and relentless and the patient too restless, a simple enucleation can be performed. A physician must, however, discriminate an isolated thrombosed external hemorrhoid from a thrombosed prolapsed hemorrhoid and should not attempt this intervention for the latter (26).

Thrombosed Hemorrhoidal Prolapse

A thrombosed hemorrhoidal prolapse usually consists of both an internal and external hemorrhoid. In case of necrosis, an emergent surgery must be performed. In this scenario, it is usually appropriate to excise the whole hemorrhoidal body including the external part. However, an emergent intervention in a hostile edematous environment exposes the patient to a significant risk of postoperative stricture. Therefore even in cases of irreducible and thrombosed hemorrhoids, if there is no necrosis, repositioning with conservative methods should be the line of treatment (27). Nevertheless, an urgent hemorrhoidectomy after a few days of relief can be a definitive treatment choice.

Crohn's Disease

A physician must always prefer conservative treatment options in patients with Crohn's Disease. Patients, who are not under steroid therapy and whose anorectal region is not affected, might be surgical candidates. Again, less invasive techniques should be preferred to excisional therapies.

Immunocompromised Patients

This patient population has an altered wound healing process. As the patients with Crohn's Disease, they are prone to surgical site infections and impaired healing. Conservative treatments must be a physician's first choice, as long as they remain feasible. If a surgery is to be performed, prophylactic antibiotics must be administered.

Pregnancy

Hemorrhoids are common during pregnancy and the symptoms usually resolve after childbirth. Normally, conservative managements with dietary cautions and sitz baths suffice to alleviate symptoms. In cases of strangulated or thrombosed hemorrhoids, a surgical intervention, preferably under local anesthesia, might be performed.

Recurrent Disease

Recurrence should be clearly distinguished from a redundant hemorrhoidal tissue. With the latter, a patient almost never experiences a sufficient relief of symptoms following surgery.

The only way to eliminate the risk of recurrence completely is to remove the CCR completely. This means after every procedure, there is a risk of recurrence. Patients can be commenced on conservative and medical treatment even after recurrence. Nevertheless, if surgery is deemed appropriate, its keypoints are as follows;

1. A ligation based technique is still feasible after a ligation based technique.
2. A ligation based technique is the treatment of choice following a conventional excisional technique, due to increased risk of stricture.
3. One should not perform stapled hemorrhoidopexy in a patient twice.

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Changes In Jaw Surgery After The Pandemic And Their Effects On Oral Health

İsmail SANCAR ŞİMŞEK¹

Introduction

The Covid-19 pandemic, which broke out in China at the end of 2019 and affected the whole world in a short time, deeply affected the production and delivery of health services, as well as all areas of life. During the pandemic process, while there were uncertainties about the exact effects of the disease, its way of spreading, which risk factors it has, and which groups it is risky for, on the other hand, there were disruptions in all areas from social life to public order, from production to education (da Silva et al, 2022; Buchy et al, 2021; Carbone et al, 2021; Sharma et al, 2021). One of the areas most affected by this situation is the areas of oral and maxillofacial surgery.

Oral health is actually an important issue that is related to many health areas and closely concerns public health and the quality of life of individuals. However, since the problems in general and non-elective health services have not been resolved, especially in developing or underdeveloped countries, oral health and related units gain importance only in the presence of a situation that affects high pain or other health conditions. (Poirier et al, 2022; Gill et al, 2022; Poudel et al, 2021; Petersen et al, 2020; Tynan et al, 2018). For this reason, it can be stated that many factors such as physicians, health workers, institutions or equipment working in the field of oral health are insufficient both compared to other health services and compared to the population.

One of the biggest challenges faced by those working in the field of oral health has been the cessation of health tourism, which has increased its orientation due to the already limited opportunities for dental health during the pandemic and especially due to very low profit margins against very high costs in the domestic market. Although health is a constitutional right of individuals and a social service, the provision of health services in reality brings a certain economic cost. In dental and oral health, this situation has a much greater effect than other health services, and external dependence is at the highest level in terms of materials (Chavda et al, 2022; Emrani et al, 2021; Imai and Tanaka, 2021; Mallah et al, 2021; Murugan et al, 2021). Moreover, this situation is valid not only for Turkey, but also for many countries in the world. Despite the dependency on foreign materials in Turkey, highly qualified workforce has made our country one of the most attractive centers in oral health and global health tourism. However, the pandemic process made this situation very difficult and closed the foreign markets where dentists and surgeons met the material-currency problems they had experienced in the domestic market.

Internally, the most important problems experienced in health services are the general perspective on oral health and the low importance given to oral health in social or cultural terms. Oral health, which was postponed or neglected even in normal times, was postponed more for many different reasons during the pandemic process and many oral health services, especially jaw surgery, were evaluated electively. Examining and revealing all these factors and the effects of the

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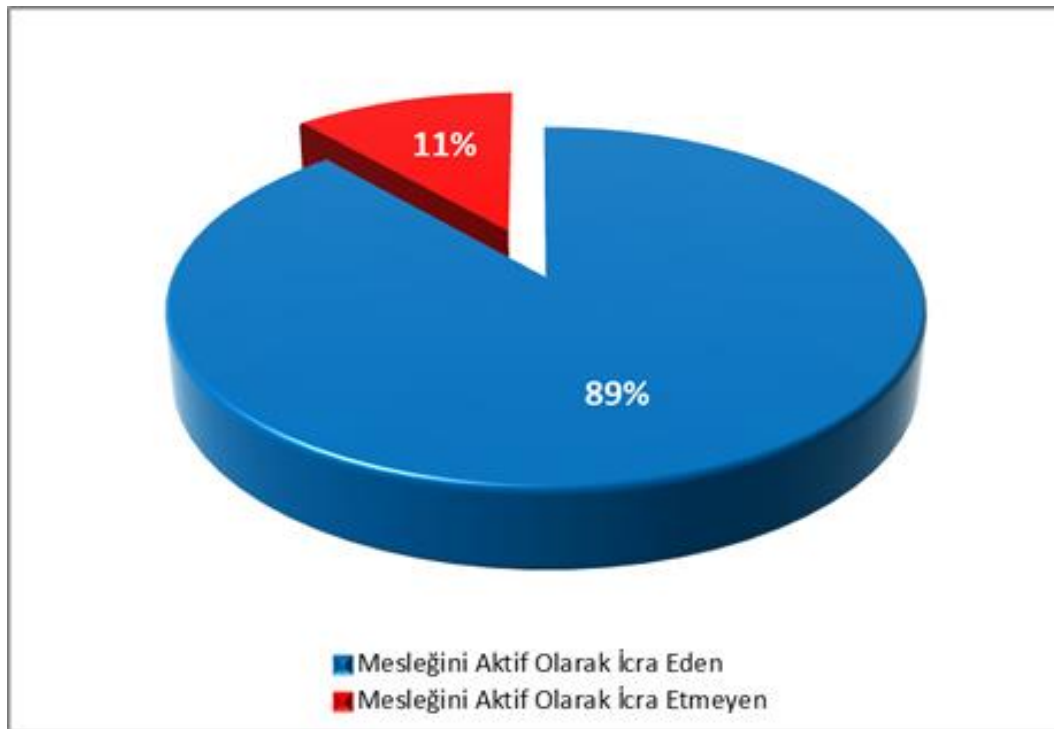
process on oral health is more important than many studies, research or clinical findings in oral health and maxillofacial surgery, and there are a series of factors that will provide a direct economic input to the field. makes a recommendation.

General Situation of Oral Health and Maxillofacial Surgery in Turkey

In Turkey, the situation of oral and maxillofacial surgery, or more generally, oral health services, should be examined under two headings as material-material and workforce. In terms of materials and materials, although production has been made for new domestic implants, consumables or coating types in recent years, these products have not yet been sufficiently accepted in the market and brands of flammable origin are dominant in the market. In this respect, it is possible to state that foreign dependency as a material in oral health is high in Turkey, therefore it is closely affected by exchange rate movements, and price stability is a disadvantage to either the dentist or the patients. In particular, the products of countries such as Switzerland and Germany, which dominate the market, almost hold the market superiority and monopoly in this field (Peker et al, 2021; Kuzu, 2019). In this respect, it can be stated that oral health in Turkey does not have a positive impression in terms of materials and materials. The increase in foreign exchange and commodity prices along with the deteriorated economies during the pandemic process, on the other hand, revealed how important domestic consumables are and how necessary a self-sufficient system is despite all globalization and transportation-communication possibilities.

The second issue is the workforce or human resources. Although there are other important workforce elements in oral and dental health, such as dental practitioners and technicians trained in associate degree departments, dentists are predominantly important in dental health services and are seen as an important indicator of the level of oral and dental health of the country. The distribution of those who became dentists in Turkey in 2021 according to their ability to practice their profession is shown in Figure 1.

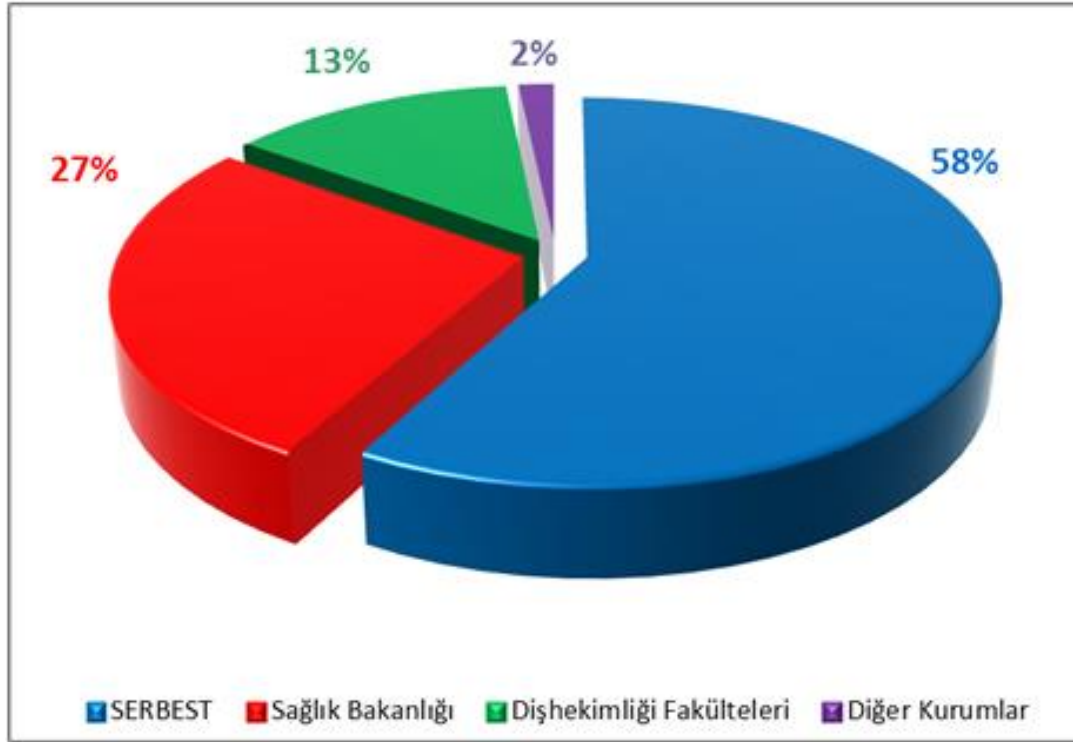
Figure 1. Distribution of those who became dentists in Turkey in 2021 according to their ability to practice their profession



Kaynak: Türkiye Diş Hekimleri Birliği, https://www.tdb.org.tr/sag_menu_goster.php?Id=92, Erişim: 22.09.2022

According to Figure 1, 89% of those who became dentists in our country in 2021 are practicing their profession, while 11% cannot. In fact, although 89% is seen as a very good figure for a profession, the rate of 11% is quite high in an area whose training constitutes a very serious economic cost according to the country's economy and the financial strength of individuals. Moreover, while the population's access to dentists in general is quite low, a significant rate of 11% cannot practice their profession either for economic or other reasons. Places where dentists work in Turkey in 2021 is shown in Figure 2.

Figure 2. Places where dentists work in Turkey in 2021



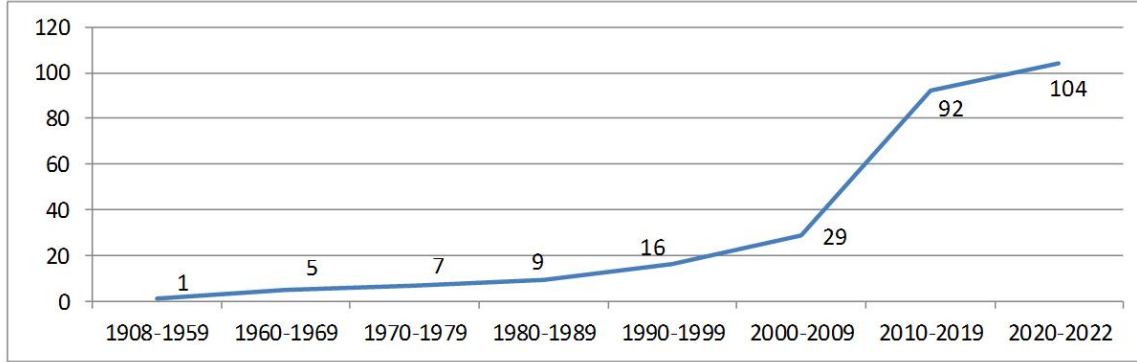
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According to the 2021 data of the Turkish Dental Association, 58% of the dentists in the country are self-employed, 27% work in the ministry of health and 13% in dentistry faculties. The ratio of those working in institutions other than these is around 2%.

In general, according to the distribution by place of work, self-employment is the most preferred type of employment in the country. Although this is followed by the ministry of health and universities, the sum of the two is not as much as the rate of self-employed. Therefore, it is possible to state that dentistry services and oral and dental health services are mostly provided in a private sector in Turkey, and this situation also brings with it costs and workload on dentists such as marketing, advertising and operation.

The change in the number of dentists in Turkey by years is given in Figure 3.

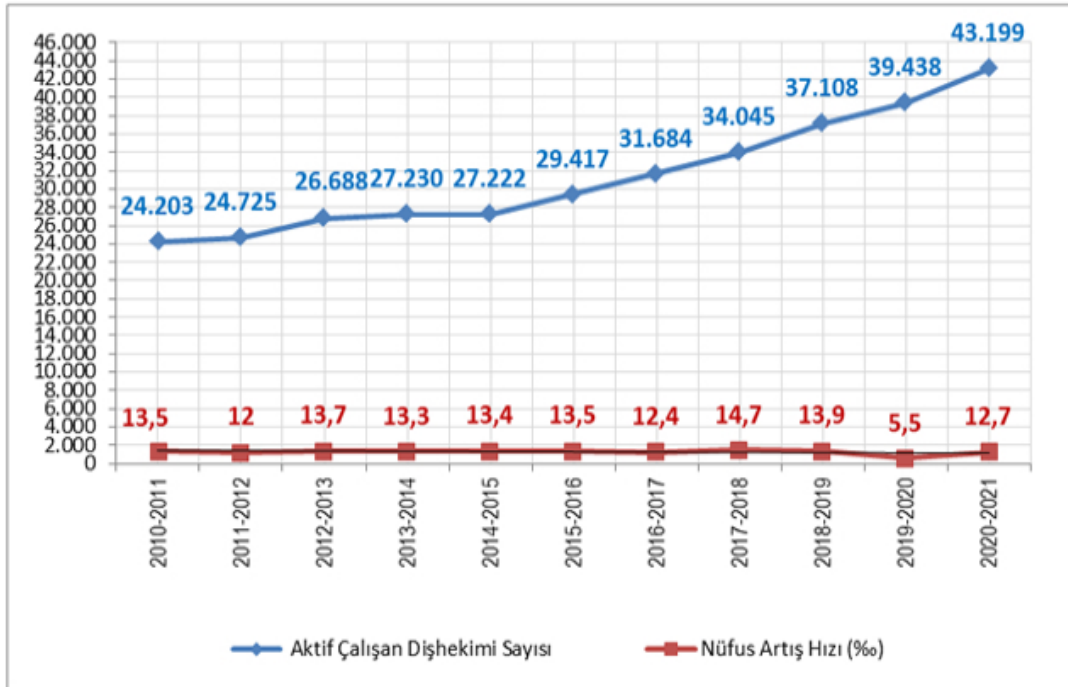
Figure 3. Change in the number of dentists in Turkey by years



Kaynak: Türkiye Diş Hekimleri Birliği, https://www.tdb.org.tr/sag_menu_goster.php?Id=84, Erişim: 22.09.2022

According to the change in the number of dentists over the years, there was a first acceleration in the 1990s and a second acceleration in the mid-2000s. Since the pandemic process in 2019, the sharp increase momentum that started in 2009 has experienced a serious decrease in the rate of increase. However, when evaluating this change, it is necessary to take into account the change in the population. The distribution of the change in the number of dentists by years according to the population is as in Figure 4.

Figure 4. Distribution of the change in the number of dentists over the years by population



Kaynak: Türkiye Diş Hekimleri Birliği, https://www.tdb.org.tr/sag_menu_goster.php?Id=387, Erişim: 22.09.2022

When the population growth rate and the number of actively working dentists are evaluated together, it is seen that the rate of dentists and the number of dentists per capita have increased significantly. It can be stated that health tourism, which has increased significantly towards the end of the 2000s, and the fact that one of the sectors that have the highest share in global health services, is oral and dental health, has a significant impact on this change.

Changes in Oral Health and Maxillofacial Surgery after the Pandemic

One of the most important changes experienced after the pandemic is the decrease in universities providing dental education. The distribution of the number of newly opened dental faculties by years, according to state and foundation universities, is given in Table 1.

Table 1. Distribution of the number of newly opened dental faculties by years, according to state and foundation universities

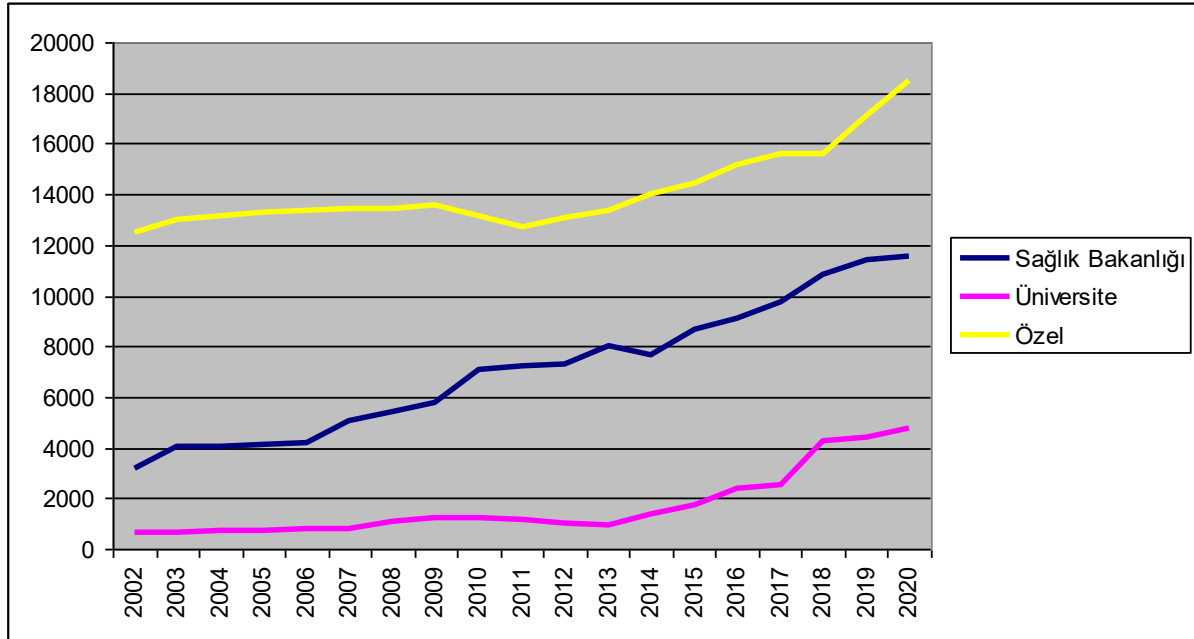
YIL	DEVLET ÜNİVERSİTESİ	VAKIF ÜNİVERSİTESİ
1908-1959	1	-
1960-1969	4	-
1970-1979	2	-
1980-1989	2	-
1990-1999	5	2
2000-2009	10	3
2010-2019	42	21
2020-2022	9	3
TOPLAM	75	29

Kaynak: Türkiye Diş Hekimleri Birliği, https://www.tdb.org.tr/sag_menu_goster.php?Id=84, Erişim: 22.09.2022

According to the data in Table 1, it is seen that the rate of increase in dentistry faculties, whose number increased rapidly between 2010 and 2019 with the effect of global health tourism, decreased after the pandemic. Although the faculties of dentistry and the process of training dentists have high economic costs and returns, due to the serious decrease in the number of international students after the pandemic, as well as the serious declines in dental health, like all other fields in global health tourism, new It is observed that the increase in the number of dental faculties has decreased significantly.

In addition to the training given at universities, there have been some changes in the employment characteristics in the sector after the pandemic. According to the 2022 research conducted by the Turkish Statistical Institute (TUIK), the distribution of the institutions where dentists work over the years is given in Figure 5.

Figure 5. According to the 2022 research conducted by the Turkish Statistical Institute (TUIK), the distribution of the institutions where dentists work depending on the years



Kaynak: TÜİK hane halkı araştırması, <https://biruni.tuik.gov.tr/medas/?kn=149&locale=tr>, Erişim: 26.09.2022

According to the data in the figure, after 2019, employment rates in both state and university decreased, while private employment increased. In fact, while the main reason for the increase in the number of people employed in the private sector after the pandemic is the decrease in the employment of dentists in public and universities, it is possible to state that there has been a serious decrease in the number of physicians going abroad due to the effect of the pandemic.

In general, the most important changes in oral health and maxillofacial surgery after the pandemic are the economic dimension of oral and dental health and the changes in the number of patients. Although health services are a constitutional and human right that individuals have in the global sense, private health services are provided in parallel with public or social health services due to insufficient or insufficient quality of public health services. Oral and dental health, on the other hand, is the area in which the private sector tends the most in terms of both the treatment and the time to reach the doctor, and the quality of the service and materials provided in public institutions. Therefore, the number of patients is of vital importance in oral and dental health services. It can be stated that the most important disruption experienced in the pandemic process and economic reasons come first in the field of oral and dental health, as in all other areas.

The situation is even worse in terms of maxillofacial surgery, and it is seen that foreign dependency in terms of materials is more effective here. In addition, serious problems have been experienced in jaw surgery due to the unlimited or insufficient interest and public support in the country, which is directed to the fields related to the pandemic. In addition to the problems of finding patients, important economic problems have been experienced due to the fact that the materials used in maxillofacial surgery are dependent on the exchange rate, the exchange rate has a sharp fluctuation, and the production has decreased all over the world due to the pandemic.

Conclusion

Although oral health is a health field that is important in terms of nutrition, good looks and acceptance in the society, it is considered within the scope of secondary health services due to the deficiencies in other health areas with high mortality rates. The company does not guarantee oral health. In fact, although the reasons for this seem to be the ones listed above, basically the most important reason for this is the cost and financial return calculations. In particular, the fact that insurance companies do not offer packages to secure dental health services within the scope of complementary health expresses the economic aspect of the situation more clearly.

At the beginning of the economic reasons, the high level of foreign dependency in terms of equipment and vehicles in the health services provided in the field of dentistry and oral health, the absence of an economic exchange rate in the country, and therefore, the foreign dependency and low predictability of each service in oral health come to the fore. Due to the unbalanced increase in exchange rates, sectors are faced with the tables that may cause harm, let alone providing income, in the ongoing oral health services. For this reason, in countries with a high level of external dependence in terms of materials and equipment, oral health services are consistently seen as elective or secondary health care services.

Turkey has overcome this negative situation or disadvantage with its qualified workforce and has become a country that provides the highest quality service at the most affordable prices in terms of oral and dental health in global markets. In this process, the low profit margin of the patients in the domestic market or the quality of health services that may be of relatively lower quality have increased as a result of the services provided by dentists and surgeons from outside. However, with the pandemic, the situation regarding oral and dental health in health tourism and global markets has been reversed. In this process, dentists and surgeons had serious economic problems, the costs, especially materials and rents, were at very high levels, therefore, these gains of the dentists, who had a very important place in the world and showed serious individual efforts, almost melted away after the pandemic. Although oral health services have increased significantly in the domestic market, it is possible to state that the main and hard blow comes from global health tourism, which has a high economic impact.

In fact, while the operations in terms of oral and dental health and especially jaw surgery continue locally in developed countries, the whole process from operations performed in jaw surgery to minor procedures has been seriously disrupted in Turkey. After the Covid-19 pandemic, even weddings, mass celebrations or ceremonies were considered obligatory and not considered elective, while even very serious procedures in jaw surgery that adversely affected health were seen as elective.

While the expected effect of the pandemic in other health fields is the formation of a health awareness and individuals to pay more attention to their health, it has brought serious economic difficulties within the framework of those who provide these services in oral and maxillofacial surgery. National action plans are needed for the continuation of the achievements of all parties working in the field of oral health, especially dentists and surgeons, with their individual efforts. The pandemic process is actually important in oral and dental health, as in any other field, in terms of revealing how negative results can be caused by external dependence in terms of materials and equipment. Again, the post-pandemic events show how fragile and in need of encouragement oral and maxillofacial surgery is economically. Oral and especially maxillofacial surgery should be strengthened and necessary support should be given, both in terms of directly affecting the health and quality of life of individuals and in terms of bringing foreign currency directly to the country from foreign markets in economic terms.

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Investigation Of Growth Differentiation Factor-15 (Gdf-15), Paraoxonase And Arylesterase Levels In The Pericardial Fluid Of Patients Undergoing Cardiac Surgery

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Introduction

The pericardium is a double-layered fibro-serous layer that covers the surface of the heart and contains the pericardial fluid. Pericardium and pericardial fluid form a closed homeostatic field that allows the heart to function easily. Interventions and illnesses may cause deterioration of the homeostatic environment and accompanying mechanical stress (1). Pericardial fluid contains growth factors, cardiac hormones and cytokines(2). Growth differentiation factor-15 (GDF-15), known as macrophage inhibitory peptide, is in the transforming growth factor β (TGF- β) superfamily. It is synthesized in response to conditions such as oxidative stress, inflammation, injury, and tissue hypoxia (3). It has been reported that it is seriously elevated in diabetes, kidney dysfunction etc.cases (4). Despite the high level of GDF-15 in patients with heart failure, the synthesis site of GDF-15 could not be fully explained because it is not seen in cardiomyocytes (5). Deaths not related to cardiovascular diseases (CVD) and CVD are associated with elevated GDF-15 levels(3). Therefore, the function of GDF-15 has not been fully elucidated (6).

Paraoxonase 1 (PON1) is a glycoprotein grouped as the aryldialkylphosphatase enzyme (7). It has anti-oxidant and anti-atherogenic effects (8). In addition, it is an enzyme that has arylesterase and lactonase functions and has the ability to hydrolyze many substrates such as glucuronides, organophosphate compounds, arylesters, and cyclic thiolactone carbonates (9). With this function, it hydrolyzes oxidized lipids from low-concentration lipoproteins (LDL) (10). Also, PON1 is associated with its anti-inflammatory and anti-oxidative properties among the subgroups of high-density lipoprotein (HDL) (8).

Examining the levels of GDF-15and antioxidant enzymes paraoxonase and arylesterase which were previously studied only at serum level in many different disease prognoses such as Pulmonary Embolism, Heart Failure, Thalassemia Major, Acute Pulmonary Proembolism, Ovarian Surface Epithelial Malignant Tumor, Left Ventricular Systolic Function Disorder, will shed light on the investigation of cardiovascular diseases.

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Material and Method

The study was initiated with the approval of the Ethics Committee numbered 16/03/38. Pericardial fluid samples were taken from the patients who underwent surgery during the opening of the pericardium after sternotomy.

Patient Population and Procedures

In this study, 40 (28 Male + 12 Female) patients who underwent open heart surgery due to various cardiovascular diseases in the Cardiovascular Surgery operating rooms of Harran University Faculty of Medicine and Mehmet Akif Inan Training and Research Hospital were included. The age of these patients was a minimum of 38 and a maximum of 85, and the mean age was 60.97 ± 8.79 years.

Average weight and height of the patients, respectively; while it is 86.23 ± 9.77 kg, 173.5 ± 4.79 cm in men, it is 73.50 ± 5.61 kg, 158.5 ± 2.16 cm in women. 97.5% of all patients had coronary bypass surgery. All of the patients reported that they smoked cigarettes. In addition, 21 of all patients had diabetes mellitus, 26 had hypertension and did not have any other diseases.

Patients with left ventricular ejection fraction $<30\%$, active infection, systemic inflammatory disease, and chronic lung and kidney disease were excluded from the study.

Obtaining Pericardial Fluid

In patients undergoing open heart surgery, after mediansternotomy with standard cardiopulmonary bypass procedures, the pericardium was opened and pericardial fluid was aspirated (at least 2 ml) with a sterile syringe. Bloody pericardial fluid samples were excluded from the study. Pericardial fluid, taken with a sterile syringe, was transferred to a sterile gel-free tube and transported to the laboratory in a cold environment. Then, the pericardial fluid in the sterile tube was centrifuged at 1000 g at $+4^\circ\text{C}$ for 20 minutes and the supernatant was taken into a sterile eppendorf tube and stored at -80°C to be studied.

Study of Pericardial Fluid Samples

Before the study, 40 samples belonging to the patients, which were removed from -80°C , were brought to room temperature and expected to dissolve. It was again passed through a second centrifuge (1000 g $+4^\circ\text{C}$ 20 min) in order to avoid the particles in it. The study was started by transferring the supernatant part on the eppendorf tube to another eppendorf tube.

Measuring the Amount of GDF 15

Elabscience Human GDF-15 (Growth Differentiation Factor 15) Eliza Kit was used for this study. This eliza kit is suitable for measuring serum, plasma and other biological fluids, and its minimum detectable dose is 14.06 pg/mL.

Paraoxonase Enzyme Activity Measurement

Paraoxonase activity, a lipophilic, hydrophobic, antioxidant enzyme linked to HDL-Cholesterol, was measured using a commercial RelAssay kit. In the method, the paraoxonase enzyme paraoxon (O,O-diethyl-O-pnitrophenylphosphate) hydrolyzes its substrate, resulting in the formation of a colored pnitrophenol product. The absorbance of the formed product was monitored in the kinetic mode at 412 nm, and the enzyme activity was expressed as U/L (11).

Arylesterase Activity Measurement

Arylesterase activity of paraoxonase enzyme, an antioxidant enzyme, was also measured using a commercial RelAssay kit. This test is based on the colorimetric measurement of phenol

released from the phenylacetate substrate by enzymatic activity. The results were expressed as kU/L because the enzyme activity was at very high levels(12).

Total Oxidant Status Level Measurement

The total oxidant status (TOS) level of the samples was measured using commercial kits. In the measurement, a colorimetric method based on the cumulative oxidation of ferrous ion to ferric ion by the oxidant molecules contained in the samples was used. Results were expressed as $\mu\text{mol H}_2\text{O}_2$ Equivalent/L (13).

Measurement of Total Antioxidant Status

Total antioxidant level (TAS) level of the samples was measured using commercial kits. The measurement method is based on the reduction of the colored ABTS* cationic radical by all antioxidant molecules in the sample, resulting in decolorization of the colored radical in proportion to the total concentrations of the antioxidant molecules. Trolox, a water-soluble analogue of vitamin E, is used as the calibrator. Results were expressed as mmol TroloxEquivalent/L (14).

Oxidative Stress Index Measurement

Oxidative Stress Index (OSI), which is shown as an indicator of Oxidative Stress, is expressed as a percentage of the ratio of Total Oxidative Status/Level (TOS) to Total Antioxidant Status/Level (TAS). When calculating the Oxidative Stress Index (OSI) of the samples, the TAS levels are multiplied by 10 to equalize the TOS levels and the units. The results were expressed as ArbitraryUnits (AU) (15).

$$\text{TOS, } \mu\text{mol H}_2\text{O}_2 \text{ Equiv. / L.}$$

$$\text{OSI} = \frac{\text{TOS, } \mu\text{mol H}_2\text{O}_2 \text{ Equiv. / L.}}{\text{TAS, } \mu\text{mol TroloxEquiv. / L.}} \times 100$$

$$\text{TAS, } \mu\text{mol TroloxEquiv. / L.}$$

Statistical analysis

Statistical analyzes were performed using the SPSS Version 17 (SPSS Inc. Chicago USA) computer program. The significance of the difference between the means of the groups was compared with the One-WayANOVA test. The relationship between the parameters was investigated by Pearson Correlation Analysis. Values less than ($p < 0.05$) were considered statistically significant.

RESULTS

GDF-15, Paraoxonase, Arylesterase, TAS, TOS and OSI values are shown in Table-I.

The lowest value of GDF-15 in the pericardial fluid of the patients was 13.57 pg/mL, the highest value was 27.45 pg/mL, and the mean was 18.21 pg/mL. Considering the paraoxonase values, the lowest value was 21 U/L, the highest value was 100 U/L, and the mean was 67.91 U/L. Arylesterase values were measured as a minimum of 321 U/L, a maximum of 399 U/L, and an average of 368 U/L.

GDF-15, Paraoxonase, Arylesterase, TAS, TOS and OSI correlation data are shown in Table-II.

When correlation analyzes were examined, a positive correlation was observed between GDF-15 and paraoxonase values ($r=0.101$), but no statistical significance was observed ($p=0.673$) (Figure-1. Correlation between GDF-15 and PON1 Levels). A negative correlation was observed

between GDF 15 and arelisterase ($r = -0.017$), no significant statistical finding was obtained ($p=0.945$) (Figure-2. Correlation between GDF-15 and Arylesterase Levels). A positive correlation was observed between paraoxonase and arelisterase ($r=0.039$), no statistically significant finding was obtained ($p=0.871$).

When the correlations of GDF-15, Arylesterase, Paraoxonase values and TAS, TOS and OSI values were evaluated, no significant statistical data could be obtained for all parameters ($P>0.05$)

DISCUSSION AND CONCLUSION

In recent studies, pericardial fluid concentrations for different cardiovascular diseases make important contributions to the detection of many pathophysiological mechanisms. Pericardial fluid contains growth factors, cardiac hormones and cytokines (2). Known as a macrophage inhibitor peptide, GDF-15 is synthesized in response to conditions such as oxidative stress, inflammation, injury, and tissue hypoxia (3).

Meloux et al. performed cerebral embolization in wistar male rats by microcirculatory intervention of the left or right internal carotid artery. They evaluated stroke lesions and cardiac function. They found that there was an increase in the levels of GDF-15 and catecholamines immediately after the stroke. They showed that GDF-15 levels were significantly increased in rats with ischemic stroke (16). In addition, they thought that GDF-15 gene polymorphism was one of the strong factors in the progression of ischemic stroke, as the stroke affected large arteries and the level of GDF-15 increased further (17)

Serum samples were obtained at each endomyocardial biopsy before and after heart transplantation by Tokavanich et al. As a result of the examinations, they observed that the level of GDF-15 increased significantly in patients with heart failure, and the level of GDF-15 decreased after transplantation. It has been stated that GDF-15 follow-up may be useful in distinguishing patients with a high risk after transplantation and may contribute to risk stratification in terms of both primary graft dysfunction and the possibility of death in these patients (18). Rothenbacher et al. investigated the relevance of baseline GDF-15 levels to overall mortality in an eight-year review of 1,470 older adults. According to the results; They explained that there is a correlation between GDF-15 levels and all factors that can cause death(19). Wollert et al. explained that, GDF-15 is associated with mortality in heart failure with atrial fibrillation. As a result of the study, it has been proven that GDF-15 levels in serum have a strong correlation with the occurrence, progression and risk prediction of CVD in the general population and patient groups(3).

HDL are high-density lipoproteins. Density, consists of subgroups that differ in terms of lipid diversity, protein content, etc. It has many functions such as prevention of inflammation, controlling the production of adhesion molecules that cause great vascular integrity, and apoptosis. It also shows anti-atherosclerotic properties by maintaining endothelial barrier activations (20). PON-1(8), which has anti-oxidant and anti-atherogenic effects, is strongly dependent on HDL (8). Sun et al. found that those without coronary artery disease (CAD) had higher arylesterase functions of PON-1 than those with CAD (21). Zuin et al. proved that PON-1 arylesterase level is decreased in patients with CAD (22). Murillo-González et al. examined serum PON-1 activity and the relationship of its polymorphisms with CVD using four different substrates and explained that it can be a guiding biomarker in the progression of CVDs(23)., Didas et al. explained that attenuation of arylesterase, an enzyme in the esterase family(24), activity may increase the likelihood of developing chronic kidney disease (CKD) in type 2 diabetes patients (T2DM) and that it can be used as a biomarker to predict the development of CKD in T2DM (25).

In our study, the level of GDF-15, paraoxonase and arylesterase, filtered into the pericardial fluid in patients undergoing open heart surgery, was determined in cardiovascular diseases and the relationship between these parameters and the pathophysiology of the disease was investigated. For this purpose, it is important that we use the pericardial fluid of patients during open heart surgery. Because, according to many studies, the concentrations of some substances in the pericardial fluid are of great importance for the diagnosis or prediction of pericardial or non-pericardial disorders (2).

Although there was no statistical difference between GDF-15 levels and arelisterase levels, a negative correlation was observed. When paraoxonase and arelisterase are examined, a positive correlation relationship is observed even though it is not statistically significant. This situation stands out as an expected situation. GDF-15 increases in conditions due to oxidative stress, on the contrary, PON-1 and arylesterase activity are observed. However, when the correlation analyzes of OSI values were evaluated in the study, the opposite results were obtained. There have been studies on GDF-15, paraoxonase and arylesterase levels in many different disease prognoses such as pulmonary embolism, heart failure, thalassemia major, acute pulmonary proembolism, surface epithelial malignant tumor of the ovary, left ventricular systolic dysfunction, only at serum level. Although there are no contradictory results and significant statistical results, the level of GDF-15, antioxidant enzymes paraoxonase and arylesterase, in the pericardial fluid, is one of the pioneering studies in the literature as far as we know, and it will be a pioneer for future studies.

Limitations of the Study

These limitations are the lack of standard methodology due to the fact that the study was conducted for the first time, the small number of patients participating in the study, the sensitive pericardial fluid taken, being adversely affected by environmental conditions, and the absence of a control group.

CONCLUSION

In our study, GDF-15, paraoxonase and arylesterase in pericardial fluid and the correlation between them were measured. As it is the first in the literature, we believe that it will shed light on future studies. For this reason, there is a need for studies with larger series, multicenter, prospective, controlled and standardized methodology regarding the parameters to be investigated in pericardial fluid.

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Table I. GDF-15, Paraoxonase, Arelisterase, TAS, TOS and OSI values ,

	Min	Max	Mean	Std. Deviation
Pericard_Gdf15	13,57	27,45	18,2170	4,17715
Paraoxonase	21,00	100,00	67,9100	19,72111
Arelisterase	321,00	399,00	368,5200	17,78363
TAS	,90	1,68	1,3340	,17901
TOS	5,70	19,00	14,5875	2,96044
OSİ	,38	1,67	1,1112	,25381

Table II. GDF 15, Paraoxonase, Arylesterase, TAS, TOS and OSI Correlation Table

	TAS	TOS	OSİ	PON1	ARE	GDF_15
TAS r	1	,11-5	-,544*	-,032	-,282	-,025
p		,628	,013	,893	,228	,918
TOS r	,115	1	,755**	-,178	-,242	-,143
p	,628		,000	,453	,303	,549
OSİ r	-,544*	,755**	1	-,104	,009	-,071
p	,013	,000		,663	,969	,767
PON1r	-,032	-,178	-,104	1	,039	,101
p	,893	,453	,663		,871	,673
AREr	-,282	-,242	,009	,039	1	-,017
p	,228	,303	,969	,871		,945
GDF_15r	,025	-,143	-,071	,101	-,017	1
p	,918	,549	,767	,673	,945	

PON1: Paraoxonase, ARE: Arelisterase, * The correlation is significant at the 0.05 level, ** The correlation is significant at the 0.01 level.

The Relationship Of Nitric Oxide With Cardiovascular Diseases And Pulmonary Hypertension

Mehmet ÖZDİN¹

Introduction

Cardiovascular diseases (CVD) are among the most important health problems affecting public health in the world. Atherosclerosis forms the basis of CVDs. Nutrition, environmental and genetic factors play an significant role in the formation. It is formed by the accumulation of fat and tissue residues under the cells lining the vein, which is the inner most layer of the vein. In this way, it progresses with stenosis or obstruction in the lumen of the vessel and causes disruption of tissue blood supply. Atherosclerosis is a multifactorial disease characterized by thickening of the vessel wall and loss of vessel elasticity due to this thickening (İlikay & et al., 2016) (Özdin, 2021a, 2021b). Atherosclerosis is the leading cause of death worldwide. One third of deaths in the world are due to these atherosclerotic diseases (Çoban, Erginel & Tuna, 2014). More than 600,000 people die annually from CVD in the United States (Mangge & et al., 2014). Various metabolic, inflammatory, infectious or hemodynamic factors are involved in the development of atherosclerosis, plaque formation and activation. Vascular endothelial dysfunction is significant in the pathogenesis of atherosclerosis causing coronary heart disease (CHD). Vascular endothelial dysfunction, which develops when endothelial cells cannot fulfill their basic functions such as vascular homeostasis and nitric oxide (NO) production, causes atherosclerotic plaque formation. Despite the complexity of CHD in its etiological, pathophysiological, clinical and epidemiological developments, it is of great importance to understand its formation mechanisms, since atherosclerosis is a preventable condition. Atherosclerotic lesions begin at an early age, aging poses a risk for CVD. Middle-aged men under 60 years of age have a 2 to 5 times greater risk of CHD than women, while the gender-dependent risk decreases with age. Family history affects 40% of the total risk profile. Other major risk factors for atherosclerosis include hypertension, type II diabetes mellitus (type II DM), elevated blood cholesterol levels, sedentary life and obesity (Özdin, 2003). (Güray, 1997).

NO was discovered in the 1970s. Robert F. Aorta drawing attention to acetylcholine, which plays a role in contraction, indicate that it is guanylate cyclase in the cytoplasm that causes relaxation. Endotel stressed that the relaxation factor is NO. It was determined to be a NO producing compound (Furchgott & Zawadski, 1990). NO is a very important molecule produced in the body by the nitric oxide synthetase (NOS) enzyme from L-arginine, a semi-essential amino acid. It must also be taken from outside. In this context, for a healthy life, attention should be paid to foods containing L-arginine and L-citrulline amino acids. The most important point to be

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considered here is to express the importance of natural NO production (Sarı, 2020). In this study, it was aimed to research the relationship of nitric oxide with CVD and pulmonary hypertension (PH).

Nitric Oxide Biosynthesis

Physiological stimuli such as acetylcholine, glutamate, bradykinin, adenosine diphosphate (ADP) activate nitric oxide synthetase (NOS) in endothelial cells. When this enzyme, which is dependent on Calcium-Calmodulin, is activated, it provides the conversion of L-Arginine and oxygen to L-Citrulline and NO by reacting. Thanks to the discovery and use of L-Arginine analogs that competitively inhibit the NOS enzyme, it has been possible to investigate the wide biological role of NO (Moncada & Higgs, 1993). NO formed as a result of this reaction diffuses into the vascular smooth muscle cells. NO activates the enzyme by fixating to the heme group of the guanylate cyclase enzyme. The activated guanylate cyclase enzyme converts guanosine triphosphate (GTP) to cyclic guanosine monophosphate (cGMP). Causes muscle relaxation in cGMP. NO, reacts with heme-containing proteins, especially hemoglobin, and is converted to nitrate (NO_3^-) and excreted in the urine (Edwards, 1995). Nitric oxide is a colorless gas and is very stable in the absence of oxygen. However, in contact with air, it quickly reacts with oxygen and turns into nitrogen dioxide (NO_2). Nitrogen Dioxide is a toxic gas that can cause tissue damage (Borland & Higginbottom, 1989).

Nitric Oxide In Cardiovascular And Pulmonary System

The main task of the vascular endothelium is to prevent the adhesion and aggregation of platelets and other blood cells. At the same time, it is to ensure the continuity of circulation by providing adequate blood flow to the blood vessels. Vascular endothelium synthesizes and secretes NO in order to exert this effect. Systemic use of NO synthesis inhibitors enhances blood pressure in small arteries and arterioles (Anggard, 1994). This indicates the importance of nitric oxide made in the vascular endothelium in regulating blood pressure and blood flow. As a matter of fact, any functional disorder in the endothelium (diabetes mellitus, homocystinemia, familial hyperlipidemia) or acquired (smoking, sedentary life, atherogenic diet) causes hypertension (Anggard, 1994, Vane, Anggard & Botting 1990). Atherosclerosis, familial hypercholesterolemia, and reduction in coronary blood flow in smokers cannot be increased by administration of acetylcholine. This is because the dilator response of endothelium to acetylcholine is decreased (Cox & et al., 1989, Celermajer & et al., 1992). Endothelium-dependent dilatation is reduced in essential hypertension and experimentally induced hypertension in animals, with clear evidence that this is due to an aberrance in the L-Arginine-NO system (Anggard, 1994). Persistent pulmonary hypertension (PPH) occurs either primary or secondary. Secondarily, it may occur due to respiratory system disorders such as perinatal asphyxia, meconium aspiration and severe respiratory distress syndrome. It may also occur due to sepsis, diaphragmatic hernia, congenital heart disease and polycythemia (Greenough, Morley & Robertson 1992). Although its exact occurrence is not known, it has been shown that increased endogenous NO activity at birth contributes to the normally observed decrease in pulmonary vascular resistance (Abman & et al., 1990). In PPH, the lack of resistance normally seen in the postnatal pulmonary vessels may be related to low endogenous NO production. In addition, the amino acid L-Arginine, which is the precursor of NO synthesis, may be insufficient in children with PPH (Vosatka, Kashyap & Botting 1994). Intravenous vasodilators, hyperventilation, high frequency oscillatory ventilation and extracorporeal membrane oxygenation are used as specific treatment in PPH. Intravenous vasodilators, prostacyclin and magnesium

sulfate treatments cannot respond to the desired level and may cause systemic hypotension ((Vosatka, Kashyap & Botting 1994, Kaapa & et al., 1985, Abu-Osba & et al., 1992). In PH, NO selectively exerts a vasodilator effect on the pulmonary vessels. Inhaled NO reduces pulmonary vascular resistance without reducing systemic vascular resistance and cannot have a vasodilatory effect on the systemic vascular bed since it is rapidly inactivated by hemoglobin as soon as it enters the bloodstream (Abman & ark., 1992, Pepke-Zabe & et al., 1991). At the same time, methemoglobin concentration rarely rises above 1-2% during inhaled NO (INO) therapy (Kinsella & Abman, 1993a). Ventilation increases perfusion compatibility in PH. In addition, it increases cardiac output, pulmonary blood flow and oxygenation. Has all the features of an ideal pulmonary vasodilator (Rossaint & et al., 1993). They can also widen the vessels in non-ventilated lung areas, disrupting the ventilation-perfusion harmony. Because of these properties, INO is used in newborns with pulmonary hypertension, in children with congenital heart disease and respiratory distress syndrome in infants. In addition, INO has been used in adults with PH, adult respiratory distress syndrome (ARDS), and bronchopulmonary dysplasia (Kinsella & Abman, 1993b).). The most popular use of NO today is inhalation in pulmonary hypertension. Pulmonary vascular diseases are a serious complication of congenital heart diseases. Hypertensive crisis in congenital heart diseases is seen particularly after open heart surgeries. PH can overrun systemic arterial pressure. As a result, the return of blood to the left atrium is reduced. Cardiac output drops rapidly (Jones, Shore & Rigloy, 1981). Reversal of PH is very difficult and hypertensive crisis has an important place in postoperative mortality and morbidity (Hopkins, & et al., 1991). Hyperoxic hyperventilation and intravenous vasodilators are widely used in the treatment of PH (Schranz, & et al., 1992). Nitric oxide inhalation is seen as an effective alternative therapy. The use of 1 ppm (parts per million) in the treatment of pulmonary hypertensive crisis gives successful results (Sellden, & et al., 1993). The use of 20-80 ppm of INO in infants with congenital heart disease who develops PH is effective in reducing pulmonary arterial pressure (Roberts, & et al., 1993). Patients with cardiorespiratory problems need to take NO at a high oxygen concentration in order to benefit from INO sufficiently. However, since toxic NO production increases when INO is given at high oxygen concentration, NO administration and monitoring should be done carefully (Foubert, & et al., 1992, Bauchet & Renaudin, 1993).

Conclusion

Although NO, which is produced in almost all healthy tissues in the organism, is structurally simple, its activity is a very complicated mediator. NO; It provides blood pressure and vascular tone balance by vasodilation by stimulating guanylate cyclase in vascular smooth muscles. It is used by inhalation in the treatment of PH in the clinic, and it is a broad-spectrum chemical that is effective on other organs.

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Phthalates and Their Toxic Effects on the Immune System

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Mehtap KARA²

Introduction

Plasticizers are added to substance and materials to increase their flexibility, workability and extensibility. Phthalates are a type of plasticizer and are also known as endocrine disrupting chemicals (EBC). Phthalates are commonly used in many products in the environment. Cosmetics, personal care products, medical devices and food packaging materials are just a few of the usage areas. The widespread and increasing use of phthalates is very worrying.

The immune system is a defense system that protects the body against external damage. It is divided into two as innate immunity (Hereditary) and acquired (Adaptive). Innate immunity gives non-specific responses while acquired immunity gives specific but slower responses. Damage to the immune system leaves the body vulnerable to pathogens.

With the increase in the use of phthalates, their toxicity on the body has come to the fore again. Studies show the toxicity of phthalates on many body systems. In addition to its endocrine disrupting properties, it has been observed that it has effects such as shortening the gestation period, increasing systolic pressure, focal testicular dysgenesis, neuronal degenerations and causing changes in immunological responses. It is thought that permanent damage remains on the baby, especially after exposure in the womb. It has been found that children are more exposed to phthalates than adults. While phthalates were used in children's toys in the past, it was noticed that children were exposed to phthalates in toys by sucking, and the use of phthalates in children playthings was banned in the following years.

When the toxicity of phthalates on the immune system was investigated, it was observed that they had positive effects on interleukin, CRP and TNF α levels. It is thought that autoimmune diseases may occur as a result of these exposures.

Exposure to phthalates is a current and important issue. Although it is not possible to completely prevent exposure due to their widespread use, its level can be reduced. Attention should be paid to the contents of the materials used, food packaging, drugs, medical devices, cosmetics and many other products. Reducing phthalate exposure is important for our health. With this compilation, it is aimed to raise people's awareness about the toxicity of phthalates to the human body and the ways of protection from phthalates.

Plasticizers

Additives that, when added to another substance, make the substance more flexible and softer are called 'Plasticizer'. This expression is mostly used for additives added to plastics (Godwin, 2017). The Council of the International Union of Pure and Applied Chemistry (IUPAC) defines plasticizer as “*A plasticizer is added to a substance and material (usually plastic or elastomer) to increase its flexibility, workability or extensibility*” (Altenhofen da Silva et al., 2011). Plasticizing technology initially

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focused on nitrocellulose type materials. In the 19th century, Alexander Parkes make a design of a new nitrocellulose plastic, using various oils to make the brittle polymer flexible. Parkes and his colleague Daniel Spill wanted to establish a plasticizer technology with this new nitrocellulose they had developed. However, they could not obtain the desired quality product with the castor oil they chose as a plasticizer (Hyatt, J.W. and Hyatt, I.S., 1870). After that, John Wesley Hyatt commercialized celluloids for the plasticizing process. In the early 1900s, phthalates and phosphates began to be used as plasticizers. Increasing interest and usage area of two important plasticizers, tricresyl phosphate and dibutyl phthalate (DBP) (Semon, W.L., 1933).

There are many classifications of plasticizers. Some of these are classifications based on chemical structures and performance. Plasticizers are divided into five according to their performance; general use, strong solvent, low temperature, low volatile and other plasticizers. Chemically, they are divided into six: phthalates, aliphatic dicarboxylic acid esters, phosphates, hydrocarbon extenders, epoxies, and others. Plasticizers can also be grouped into primary plasticizers and secondary plasticizers. Primary plasticizers are substances that are present in the plasticization system as the sole plasticizer or a high level of plasticizer. Secondary plasticizers are used in combination with primary plasticizers to improve the plasticization process, improve certain performances such as reduced flammability, or reduce costs (Godwin and Krauskopf, 2007).

There are different theories explaining the plasticization mechanism. Some of these are lubricant, gel, and free volume theories. According to the lubricant theory, the plasticizer acts as a molecular lubricant that allows the free move of polymer chains over each other if a force is applied to the polymer. According to the gel theory, the polymer molecules are loosely attached to each other at different points. The plasticizer added to the polymer increases the movement of the polymer chain (Daniels, 2009). According to the free volume theory, the free volume represents the internal volume in the polymer. As the free volume increases, the movement of the polymer chains will be easier, and thus the flexibility will increase (Cadogan, D.F. and Howick, C.J., 2000).

Plasticizer Types

Aside from having a lot of areas of use, plasticizers also have a lot of types. Some of their name and consumption rates in 2014 are given in Table 1. (Godwin, A., 2017).

Table 1. : Plasticizer types and their consumption rates in 2014 (Godwin, A., 2017).

Plasticizer Type	Consumption Rate (%)
Phthalate Esters	75
Terephthalate	7
Dibasic acid esters	3
Epoxy plasticizers	3
Trimellitate esters	2
Benzoates	2
Cyclohexanoates	<2
Polymeric Plasticizers	<2
Phosphates	1
Citrates	<1
Others	3

Annotation: The classification of plasticizers according to their chemical types and their consumption percentages in 2014 are given in Table 1. Compared to today, the biggest change in this table was seen in terephthalate plasticizers. With the prohibition of the use of some phthalate esters in children's toys, an

increase in the use of terephthalate has been observed and it is predicted that this increase will continue in the future (Bolton, D., 2013).

Immune System

The Immune System is body defense system that protects the body against diseases, foreign substances that may come from outside, and their pathogenic effects. The main function of the immune system is to recognize foreign substances and develop the necessary defense mechanisms against them (Songu and Katmış, 2012).

The immune system can be classified as innate and the adaptive immune system. Innate immunity is a non-specific defense system that is formed according to the innate characteristics of the person. It cannot distinguish the difference of pathogens. Adaptive immunity, on the other hand, is in a state of change. It responds specifically by distinguishing foreign substances and pathogens from its own cells (Songu and Katmış, 2012). The differences between the adaptive immunity and the innate immunity are given in Table 2. (Tomar, N and Rajat, K.D., 2014).

Table 2.: Plasticizer types and their consumption rates in 2014 (Godwin, A., 2017).

The Innate Immune System	Adaptive Immune System
Non-specific response	Specific response
Immediate response	Delay between antigen exposure and response
No immunological memory	Retains immunological memory
Found in nearly all forms of life	Found only in jawed vertebrates

Annotation: The innate immune system cannot distinguish between pathogens. In contrast, the adaptive immune system specifically distinguishes pathogens but responds more slowly than innate immunity. While there is immunological memory originating from T memory cells in adaptive immunity, these cells are not found in innate immunity.

Lymphocytes, an important member of the adaptive immunity, contribute to cellular and humoral immunity. In cellular immunity, the effect of T lymphocytes and macrophages is dominant. Macrophage, one of the most effective immune cells, is activated after encountering T cell cytokines (Songu and Katmış, 2012). The innate immune system is called by this name because it provides protection against pathogens without being dependent on an external factor. The innate immune system reacts immediately when it encounters pathogens (McComb, S. et al., 2019). The main four defense barrier types of innate immunity are; anatomical (skin, mucosa), physiological (temperature changes, pH degree), phagocytic cells as monocytes, neutrophils, tissue macrophages and inflammatory (serum proteins). Cells of the innate immunity are; phagocytes, neutrophils, macrophages, natural killer (NK) cells, mast cells, basophils, dendritic cells and eosinophils. The classified version of the immune system is given in Figure 1. (Tomar, N and Rajat, K.D., 2014). Antibodies are an element of humoral immunity together with cytokines and complement system. Immune system cells can be found in the body and also have a network of organs. It is distributed throughout the body in a way that can quickly stop the spreading infection. The innate immune cells are usually formed in the bone marrow and reside in the blood and tissues. The cells of the adaptive immune system; T and B lymphocytes, mature in the thymus and bone marrow, also known as primary immune organs, and then remain in the lymphatic tissue. The accommodation sites of these lymphocytes are known as secondary immune organs. Some of these organs are; lymph nodes, spleen, appendix, tonsils, mucosa-associated lymphoid tissues, and adenoids (McComb, S. et al., 2019).

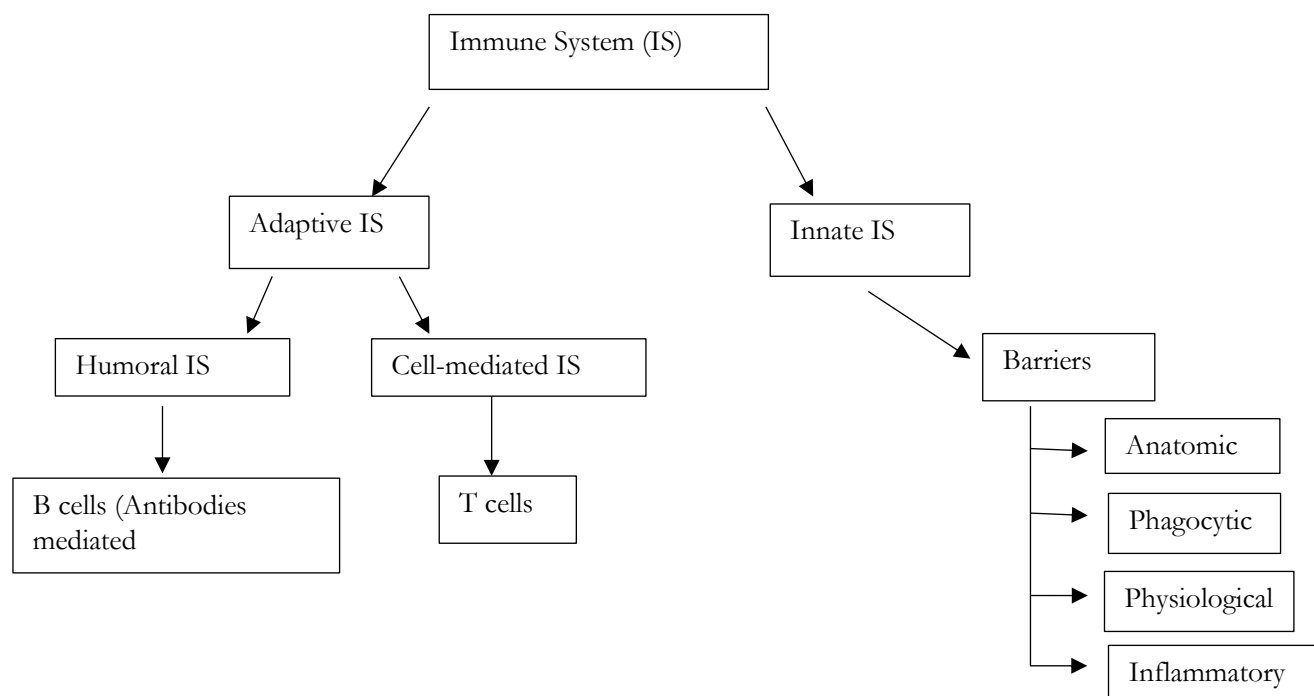


Figure 1.: Types of the Immune System (Tomar, N. and Rajat, K.D., 2014)

Phthalates

Plastics have brought enormous benefits to society since they were invented, but they have also had negative effects on the environment and human health. People are constantly exposed to plastics. Of these, the most worrying are; Phthalates are known as endocrine disrupting chemicals (EBK). Phthalates are mainly used for their softening effects in polyvinyl chloride (PVC) plastics (Wang, Y. and Qian, H., 2021). Phthalates also act as lubricants, solvents and impart positive effects to the products they are added to. There are potential routes of exposure by ingestion, inhalation, intravenous injection and skin contact (Schettler, T., 2006). While phthalate consumption is increasing, more than 3000000 metric tons of phthalates are used worldwide every year (Lyche, J.L., et al., 2009).

Phthalates are diester structured containing a benzene ring with two ester functional groups. They have low water solubility, with increasing chain length and high molecular weight. Therefore, while low molecular weight phthalates such as DMP (Dimethyl phthalate) and DBP are more soluble in water, phthalates with high molecular weight and long alkyl chains have low water solubility (Lyche, J.L., et al., 2009). Phthalates are generally lipophilic and this property affects their spreading and environmental degradation properties (Schettler, T., 2006). The general chemical structures of phthalates are shown in Figure 2 (EPA, 2012).

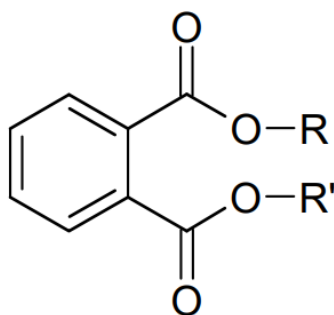


Figure 2.: General chemical structures of phthalates (EPA, 2012).

Annotation: Dialkyl ortho-phthalates (or phthalate esters) have the chemical structure shown above. R groups can be linear, branched, linear-branched or circular (EPA, 2012).

Phthalate Types

Phthalates are phthalic acid diesters and they are classified depends on molecular weight as high and low. High molecular weight phthalates include various compounds that are widely used to more flexible and durable plastics. Among them, the most widely used chemical is diethylhexyl phthalate (DEHP) (Filardi, T. et al., 2020). Phthalates such as DEHP, DBP, diethyl phthalate (DEP), di-isononyl phthalate (DiNP) and di-iso-decyl phthalate (DiDP) are mainly used to produce PVC (Wang, Y. and Qian, H. , 2021). Low molecular weight phthalates are common mainly in personal care products, cosmetics, pesticides and food packaging plastics (Filardi, T. et al., 2020). With low molecular weight short-branched phthalates such as DMP and DEP are widely produced and used in medical products and medical devices. Phthalate types and where they can be found are shown in Table 3. (Wang, Y. and Qian, H., 2021).

Table 2.: Phthalate Types and where they can be found (Wang, Y. and Qian, H., 2021)

Compounds	Abbreviation	Where It Can Be Found
Diethylhexyl phthalate	DEHP	Plasticizer
Dibutyl phthalate	DBP	Nail polishers; plasticizer; an additive to adhesives or printing inks
Diethyl phthalate	DEP	Toothbrushes; automobile parts; tools; toys; food packaging; cosmetics; insecticides; aspirin
Mono-(2- ethylhexyl) phthalate	MEHP	Vinyl tiles; food conveyor belts; carpet tile; artificial leather

Annotation: It is almost impossible to avoid exposure to phthalates, as they are ubiquitous in the environment. In addition, exposure to phthalates is increasing due to the use of plastic. Some of the phthalates we encounter in our daily lives and they can be found are given in the table above (Wang, Y. and Qian, H., 2021).

Phthalate firstly metabolized with hydrolysis after absorption into the cell, then conjugated with hydrophilic glucuronide conjugate and catalyzed via uridine 5'-diphosphoglucuronyl transferase. The type of phthalates exert their toxicological effect in the body with different mechanisms. Short-branched phthalates hydrolyzed into monoester phthalates and then excreted via urine, and long-branched phthalates metabolized as hydroxylation and oxidation, which are then excreted in urine and feces (Wang, Y. and Qian, H., 2021).

Toxic effects of Phthalates

In 2006, the National Toxicology Program (NTP) classified the effect of DEHP on the reproductive system development of one-year-old boys as 'concerning'. At the same time, the effects on the reproductive system of boys older than one year of age and exposed in the womb have been classified as 'somewhat alarming'. Some studies show that exposure to DEHP increases the risks of asthma, bronchial obstruction, ADHD and its symptoms and altered pregnancy durations. The National Research Council concluded that multiple phthalates can cause cumulative adverse effects on male reproductive system development (EPA (United States Environmental Protection Agency), 2017).

Scientific Committee on Toxicity, Ecotoxicity and the Environment has reached tolerable daily intake values (TDI) for phthalates based on NOAEL (No Observable Adverse Effect Level) levels from animal studies. They concluded that predominantly used phthalates such as DEHP and DiNP had the lowest calculated TDI values (Heudorf, U. et al., 2007). The NOAEL and TDI values of phthalates in toys are shown in Table 3. (CSTEE, 1998)

Table 3.: The NOAEL and TDI values of phthalates in toys (CSTEE, 1998).

Phthalate	NOAEL value mg/kg/day	Tolerable daily intake µg/kg/day
DiNP	15	150
DEHP	3,7	37
DNOP	37	370
DiDP	25	250

Phthalate esters and their metabolites residues commonly detected in different consumer products, urine, breast milk, and amniotic fluid. Due to phthalates can also cross the placenta, they could be teratogenic. Babies and young children consume more calories and relatively higher levels of fatty foods than adults. Children's estimated total dietary DEHP intake is highest, compared to younger adolescents. In addition to dietary exposure, oral phthalate ingestion also occurs via mouth, suck, or chew toys. The passage of DEHP from toys with saliva solubilization. With the release from some toys being higher than the maximum pass-through level determined by the Scientific Committee on Toxicity, Ecotoxicity and Environment, the European Union (EU) banned the use of phthalate types in children's toys and childcare products in 1999. In 2005 the use of DEHP banned (Lyche, J.L., et al., 2009). In the United States (USA), the use of DiNP is still allowed in toys, and it has been shown that children take up significant levels of DiNP as a result of chewing activities (Kavlock et al., 2002). Human exposure can either be assessed based on concentrations found in environmental samples and food, or indirectly reconstructed from urinary metabolite levels that reflect internal exposure. There is substantial evidence to suggest that phthalate types negatively effects the male reproductive system functions, and it has therefore been suggested that they may have a similar mechanism of action. High doses phthalates inhibit testosterone production in both fetal and adult Leydig cells. In contrast, low doses of phthalates

increase the number of adult Leydig cells and increase testosterone production. Side effects of phthalate exposure related with dose and exposure duration. Serum thyroid hormone (TH) and thyroid stimulating hormone (TSH) levels disruptions were related with MEHP concentrations that can be indirectly affects the male and female reproductive system. Estimated concentrations of DEHP in the general population were found to be highest in children and decrease with age. The critical period for effects on male reproductive development appears to be late pregnancy and the postpartum period. Recent data have shown that phthalates with a similar mechanism of action, when administered as a mixture, cause cumulative, dose-added effects on fetal testosterone production and testicular histopathology (Lyche, J.L., et al., 2009).

There are many studies showing that phthalates have serious toxic effects on many systems such as cardiovascular system, reproductive system, nervous system and endocrine system. (Mariana, M. & Cairrao, E., 2020) (Holahan, M.R., & Smith, C.A., 2015) (Hannon, P.R. & Flaws, J.A., 2015) (Wang, Y. et al., 2019) (Holahan, M.R. and Smith, C.A., 2015) (Philippat, C. et al., 2012).

According to a study, deodorants, perfumes, shampoos, creams, make-up materials, soaps and lotions are hygiene and beauty products that are used by pregnant women and contain high levels of phthalates. DEHP is the most abundant phthalate in make-up products. However, this rate may vary according to factors such as geographical conditions, diet, plastic use (Bustamante-Montes, L.P., 2021). Exposure to phthalates during a sensitive period such as pregnancy causes metabolic dysfunction in the mother and disrupts fetal development. (Filardi, T. et al., 2020).

Phthalates and Their Toxic Effects on the Immune System

For a functioning immune system, organs, cells and secretions provide body homeostasis. The first line of defense is innate immune reactions. Stimulation of this system depends on previous encounter with the pathogen; therefore, some time is required for it to react (Toskala, E., 2014). Epidemiological studies have investigated the possible relationship between exposure to PVC or phthalates and respiratory tract diseases in environments such as the home and workplace (Jaakkola, J.J. and Knight, T.L., 2008). It has been suggested in both in vivo and in vitro experimental studies that phthalates have adjuvant effects on immunoglobulin production, enable differentiation from T helper cells, affect cytokine secretion, increase enzyme and histamine secretion, and increase the phagocytic ability of innate immune cells (Bornehag, C.G. and Nanberg, E., 2010). Monocytes and macrophages are cells of the innate immune system. Monocytes are found between the bone marrow and the bloodstream. They can migrate between tissues and differentiate into macrophages in the tissues they pass through (Verschoor, C.P. and Puchta, A., 2012).

In studies investigating the relationship between inflammatory responses and phthalate esters; while there correlation between high molecular weight phthalate esters and CRP (C-reactive protein), IL-6 (Interleukin-6) and IL-10 (Interleukin-10), no relationship was found with low molecular weight phthalate esters. According to studies, phthalate ester metabolites MBzP (monobenzyl phthalate), MiBP (monoisobutyl phthalate) and MnBP (mono-n-butyl phthalate) are directly associated with high levels of CRP, while other phthalate ester metabolites MCNP (monocarboxynonyl phthalate) and MCP (mono- 3-carboxypropyl phthalate) is related with high levels of IL-6 (Franken, C., et al., 2017). Studies with DEHP also show that phthalates may cause dysregulation in inflammatory responses and increase TNF α (Tumor necrosis factor) production from monocytes and macrophages Hansen, J.F., et al., 2015a). In another review, the effects of two phthalate ester metabolites MEP and MiBP on inflammation elements (TNF α , CRP, alkaline phosphatase (ALP)) and absolute neutrophil count, ferritin, fibrinogen, IL-1 β , IL-6 and IL-10 were investigated. The results found were not in line with other studies (Sweeney, M.R. et al., 2019). In addition, phthalate esters were found to be positively associated with the anti-inflammatory cytokine IL-10. The effect of phthalates on both pro-inflammatory and anti-inflammatory responses has also been demonstrated by two different in vitro studies (Hansen, J.F., et al., 2015b).

Both studies show an increase in IL-10 levels after exposure to phthalate esters. IL-10, produced by natural killer (NK) cells during a systemic infection, acts as regulators that stabilize inflammation and limit immune-mediated damage to the host (Martinez-Espinosa, I., et al., 2021). IL-10 secreted by NK can also inhibit the function of effector T cells and reduce the efficacy of anti-tumor immunity (Sun., H., et al., 2021). Increasing IL-10 levels reduces inflammation-mediated damage but can also lead to weak immune responses against the tumor. As mentioned before, phthalates are divided into two according to their molecular weight. High molecular weight phthalates and low molecular weight phthalates differ in terms of industrial use and exposure routes (North, M.L., et al., 2014). While low molecular weight phthalates are used in personal care products, medicines and other consumer products; high molecular weight phthalates are mostly found in PVC food packaging, home furniture and other construction materials (Beszterda, M. and Franski, R., 2018).

It is thought that DEHP and monoethylhexyl phthalate reduce the permeability of the mitochondrial membrane in the human lymphoblast cell line (TK6), thus leading to Reactive oxygen species (ROS) production and caspase activation (Rosado-Berrios, C.A. et al., 2011) (Nowak, K. et al., 2019).

Discussion

Phthalates are a group of plasticizers added to plastics to give them flexibility and reduce their brittleness. They are also called endocrine disrupting chemicals. According to their molecular weights; they are divided into two groups as high molecular weight phthalates and low molecular weight phthalates. They are found in many diverse and different areas such as food packaging, cosmetics, medical devices, drugs, construction materials, personal care products. Exposure is inevitable as these chemicals are ubiquitous in the environment.

The immune system is a system that protects the body against pathogens that may come from outside and develops a defense system. There are two forms of immunity, innate and adaptive. Innate immunity produces non-specific responses and is the first line of defense. Elements of the innate immune system are natural killer cells, macrophages, dendritic cells, and skin. Adaptive immunity, unlike innate immunity, responds specifically and is in constant change. Adaptive immunity elements are T and B lymphocytes. It is divided into humoral and cellular immunity. Antibodies, which are an element of the humoral system, form an important part of the immune system. Antibodies recognize the part of the pathogen called the antigen. The system called the complement system is involved in the formation of the response in both the acquired immunity and the innate immunity. There are three types of pathways that activate the complement system; the classical pathway, the alternative pathway, and the mannose-binding lectin pathway.

The toxic effects of phthalates on humans are classified as worrisome. Many studies have investigated the toxic effects of phthalates on the systems of the human body. It has been observed that changes occur in the time of birth as a result of exposure during pregnancy. It is known to reduce testosterone production and delay puberty when exposed to high doses. Exposure is very important especially in the womb and in the pediatric period.

Studies show that phthalates have positive effects on interleukin levels, CRP and TNF α . It is also thought that phthalate exposure may lead to autoimmune diseases.

Exposure to phthalates is an important risk factor today. In order to protect ourselves from phthalates, we should pay attention to the packaging materials of the materials we use and the food we eat. Since there is a possibility that phthalates can be transmitted through the air, we should ventilate the environments we are in regularly. Even if it is not possible to be completely protected from phthalates, which are ubiquitous in the environment, it is important for our health to minimize exposure.

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Can Neutrophil-to-Lymphocyte Ratio and Platelet-to-Lymphocyte Ratio Be Used to Predict Viral Etiology in Patients with Lower Respiratory Tract Infection in the Pediatric Intensive Care Unit?

Merve MISIRLIOGLU¹

Introduction

Viral infections are the most common causes of lower respiratory tract infections (LRTIs) in childhood. Failure to determine the etiology in patients with LRTIs in the pediatric intensive care unit (PICU) may lead to inappropriate treatment, unnecessary antibiotic use, and development of antibiotic resistance, and prolonged hospitalization and intensive care unit stays (1-2).

The neutrophil/lymphocyte ratio (NLR) is calculated using the neutrophil and lymphocyte values in the complete blood count and is an easily accessible, new biomarker that has been gaining in popularity recently and can predict the progression and mortality in many rheumatologic, cardiac and endocrinologic inflammatory diseases. It is thought to indicate the severity of inflammation. Platelet/lymphocyte ratio (PLR) is used to predict the inflammatory process, disease activity, response to treatment and prognosis in inflammatory diseases and malignancies (2).

Identification of the cause of the disease; It ensures that unnecessary treatment agents are not used, the clinical course is predicted, and necessary treatment manipulations and precautions are taken. In this study, it was aimed to investigate NLR and PLR values in predicting the etiology of viral respiratory pathogens in patients with LRTIs admitted to the PICU due to respiratory failure.

Materials and Methods

Samsun Training and Research Hospital pediatric intensive care unit provides service with 6 beds as a secondary level. Between October 2021 and April 2022, 126 patients were hospitalized in the PICU. 26 patients who were admitted to the PICU due to respiratory failure and whose viral panel was studied from nasopharyngeal swab were included in the study. Patient data were obtained retrospectively from patient files in electronic media. Patients' age, gender, month of illness, length of stay in hospital and PICU, NLR, PLR values with leukocyte, lymphocyte, neutrophil and platelet measurements in complete blood count, respiratory failure treatment approaches (high flow nasal cannula oxygen therapy, noninvasive and invasive mechanical ventilation) was recorded.

Patients in the 1 month-18 age group who were hospitalized due to respiratory failure due to lower respiratory tract infection in the PICU and whose respiratory tract viral panel was studied from nasopharyngeal swab were included. Patients with respiratory failure other than lower

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respiratory tract infection, patients under the age of 1 month and over 18 years of age, those hospitalized outside of the PICU, and patients whose respiratory tract viral panel was not studied from nasopharyngeal swab were excluded from the study. In addition, cases with conditions that may affect the hematological parameters of the patients (chemotherapy, hematological malignancy, immunosuppression) were not included in the study. Approval was obtained from Samsun University Clinical Research Ethics Committee for the study (SUKAEK-2022/7/6).

Statistical analysis

SPSS version 21.0 (IBM SPSS) program was used for statistical analysis. Risk factors affecting respiratory viral panel positivity were analyzed by Binary Logistic Regression Analysis and Enter method was used to include parameters in the model. ROC Analysis was used to examine the parameters determining respiratory tract infection positivity, and DeLong Test was used to determine which of the parameters was better in determining positivity. Analysis results were presented as mean $\pm$ standard deviation. Significance level was accepted as $p < 0.05$.

Results

In the pediatric intensive care unit, 126 patients were hospitalized for a period of 7 months. 26 patients who were admitted to the PICU due to respiratory failure due to lower respiratory tract infection and whose viral panel was studied from nasopharyngeal swab were included in the study. 61.50% ($n=16$) of the patients were male and their mean age was 20.77 ± 39.50 (1-204) months; respiratory viral pathogen was positive in 80.76% ($n=21$). In 5 patients, dual pathogens were detected and co-infection was present. There was no mortality in any of our cases. The length of stay in the PICU was 4.88 ± 2.97 (3-14) days and the length of stay in the hospital was 9.15 ± 3.25 (4-19) days. The characteristics of the patients are given in Table-1.

The results of whole blood parameters checked during the patients' admission to the intensive care unit; It is shown in Table-2. NLR was 3.18 ± 3.65 (0.19-12.14) and PLR was 142.42 ± 108.62 (12.48-350.90). To determine the parameters that can be used to predict respiratory viral panel positivity; the effects on positivity were examined by Binary Logistic Regression Analysis. univariate and multivariate models were established. Since the PLR and NLR parameters cannot enter the model at the same time (multi-connection problem), two multivariate models have been established in which both are included separately. Other parameters were also excluded from the model due to the multicollinearity problem. As a result of the analyzes performed, complete blood count, biochemistry and blood gas parameters were not found to be an effective risk factor for respiratory tract viral panel positivity ($p > 0.05$) (Table-3).

Discussion

Lower respiratory tract infections; pneumonia and bronchiolitis consist of bronchitis. However, since it is difficult to distinguish from each other, especially in pediatric patients in the younger age group, this is the definition that covers these tables (3). Although the agents of infection vary according to age, viruses appear as the most common pathogens under the age of five (4). Respiratory Syncytial Virus (RSV) is the most common cause of bronchiolitis and pneumonia, and is a virus that causes epidemics especially in winter and spring (5). In our study, the most common agents were RSV and Rhinovirus. In another study conducted in critically ill children in our country, RSV was found to be the most common, and Rhinovirus was the second most common (1).

As a result of the differentiation of stem cells in the bone marrow, leukocytes, erythrocytes and platelets are produced. Although the density of the leukocyte series varies according to age, the majority of them are neutrophils. In the case of inflammation, the lymphocyte count decreases, while the neutrophil and platelet count increase. It is thought that NLR indicates the severity of inflammation (6). Numerous studies conducted in pediatric patient groups have reported an association between increased NLR and PLR and inflammation. NLR and PLR can be easily used in the diagnosis of diseases with systemic inflammation such as Henoch-Schoenlein Purpura, asthma, acute pyelonephritis, and acute appendicitis and in monitoring their progression. However, NLR was found to be a factor affecting mortality and morbidity in intensive care patients (7-9). In an adult study, it was reported that NLR could be a useful biomarker for mortality and acute respiratory distress syndrome (ARDS) development in patients with miliary tuberculosis and ARDS (10). Another hematological parameter whose number changes in inflammations is platelets. The platelet count increases not only in case of acute inflammations, but also in chronic inflammatory diseases. There are data showing that PLR, which can be used to predict disease activity, response to treatment, and prognosis in the inflammatory process, affects mortality and morbidity (11,12).

In a PubMed-based study we conducted, we could not find any publication in the literature examining the relationship between lower respiratory tract infection etiology and NLR and PLR in the pediatric age group. To the best of our knowledge, our study is the first study on the relationship between NLR and PLR in patients hospitalized in the pediatric intensive care unit due to respiratory failure due to LRTIs. In a study conducted in adult COVID-19 patients, they stated that there was a statistically significant difference between NLR healthy and covid patients, who also received and did not receive intensive care support (13). In a study conducted with 216 adult pneumonia patients in the intensive care unit, NLR and PLR were studied in the differentiation of bacterial and viral pneumonia, but no difference was found; stated that it is not an ideal biomarker for viral / bacterial differentiation (14). In our study, no statistically significant difference was found in NLR and PLR values in predicting respiratory viral positivity.

As a result; In respiratory tract infections, the correct diagnosis and the application of the right treatment can reduce the use of unnecessary antibiotic or antiviral treatment, prevent drug resistance and prevent morbidities that may develop. PLR and NLR are easily accessible and inexpensive parameters that can be used to detect the inflammatory state and do not require complex calculations. According to the results of our study, we determined that PLR and NLR could not be used to predict viral etiology in critically ill children with LRTIs; however, prospective studies with larger patient groups are needed.

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Tables

Table 1. Demographic and clinical characteristics of the patients

Patients' characteristics	n (%)
Age Groups 1 month-3 months: 3 months-5 years: ≥5 years:	6 (23.1%) (2 patients positive / 4 patients negative) 18 (69.2%) (17 patients positive / 1 patient negative) 2 (7.7%) (2 patients positive)
Distribution by month of follow-up October: November: December: January: February: March: April:	3 (11.6%) (2 patients positive / 1 patient negative) 8 (30.7%) (6 patients positive / 2 patients negative) 2 (7.7%) (2 patients positive) 1 (3.8%) (1 patients positive) 2 (7.7%) (1 patients positive / 1 patient negative) 4 (15.4%) (4 patients positive) 6 (23.1%) (5 patients positive / 1 patient negative)
Detected Viral Pathogens Rhinovirus: H.bocavirus: RSV A/B: Enterovirus: Parainfluenza 4: Coronavirus OC43: H.metapneumovirus: Parechovirus: Parainfluenza 3:	7 (26.9%) 2 (7.7%) 7 (26.9%) 2 (7.7%) 2 (7.7%) 2 (7.7%) 2 (7.7%) 1 (3.8%) 1 (3.8%)
5 patients with co-infection 1. patient: 2. patient: 3. patient: 4. patient: 5. patient:	Rhinovirus + H.bocavirus Rhinovirus + Enterovirus Rhinovirus + Parainfluenza 4 Rhinovirus + Enterovirus RSV A/B + Coronavirus OC43
Co morbid Disease	Yes: 7(26.9%) None: 19 (73.1%)
Individual with respiratory tract infection at home	Yes: 13(50.0%) None: 13(50.0%)
History of recurrent respiratory infection	Yes: 7(26.9%) None: 19 (73.1%)
History of previous inhaler treatment	Yes: 10 (38.5%) None: 16 (61.5%)
High flow nasal cannula oxygen	Yes:25 (96.2%) None:1 (3.8%)
Noninvasive mechanical ventilation	Yes: 3 (11.5%) None:23 (88.5%)
Invasive mechanical ventilation	Yes: 3 (11.5%) None: 23 (88.5%)

Table 2. Results of whole blood parameters checked during the patients' admission to the pediatric intensive care unit

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	Mean $\pm$ SD Median (minimum-Maksimum)
WBC $10^9/L$	12.05 $\pm$ 4.72 11.04 (4.28-23.35)
ANC $10^9/L$	6.66 $\pm$ 4.66 5.55 (1.40-20.77)
ALC $10^9/L$	4.18 $\pm$ 3.11 3.75 (0.90-13.30)
PLT $10^9/L$	331.50 $\pm$ 117.12 332.00 (152.00-582.00)
NLR	3.18 $\pm$ 3.65 1.38 (0.19-12.14)
PLR	142.42 $\pm$ 108.62 107.08 (12.48-350.90)
HGB gr/dL	10.93 $\pm$ 1.74 10.85 (8.4-15.6)
HCT %	32.22 $\pm$ 5.58 32.30 (23.9-44.7)

WBC: White blood cell, ANC: Absolute neutrophil count, ALC: Absolute lymphocyte count, NLR: Neutrophile-to-lymphocyte ratio, PLR: Platelet-to-lymphocyte ratio, HGB: Hemoglobin, HCT: Hematocrit.

Table 3. Examination of risk factors affecting respiratory tract viral panel positivity

	Viral Panel		Univariate		Multivariate 1		Multivariate 2	
	Negative	Positive	OR (%95 CI)	P value	OR (%95 CI)	P-value	OR (%95 CI)	P-value
WBC	11.83 $\pm$ 2.91	12.1 $\pm$ 5.14	1.013 (0.818 - 1.254)	0.907	---	---	---	---
ANS	6.09 $\pm$ 3.47	6.8 $\pm$ 4.96	1.037 (0.825 - 1.305)	0.754	---	---	---	---
ALS	4.33 $\pm$ 1.84	4.14 $\pm$ 3.38	0.981 (0.718 - 1.339)	0.903	---	---	---	---
PLT	402 $\pm$ 108.52	314.71 $\pm$ 115.12	0.993 (0.984 - 1.002)	0.148	---	---	---	---
NLR	2.08 $\pm$ 2.37	3.44 $\pm$ 3.9	1.141 (0.803 - 1.62)	0.461	---	---	0.953 (0.505 - 1.798)	0.882
PLR	140.73 $\pm$ 115.57	142.82 $\pm$ 109.89	1 (0.991 - 1.009)	0.968	0.993 (0.971 - 1.016)	0.556	---	---
HGB	9.82 $\pm$ 1.66	11.2 $\pm$ 1.69	1.895 (0.848 - 4.235)	0.119	0.876 (0.089 - 8.609)	0.910	1.119 (0.095 - 13.154)	0.929
HCT	28.78 $\pm$ 6.63	33.04 $\pm$ 5.14	1.198 (0.944 - 1.522)	0.138	---	---	---	---
MPV	8.18 $\pm$ 1.04	8.61 $\pm$ 1.19	1.495 (0.522 - 4.278)	0.454	3.699 (0.227 - 60.23)	0.358	3.526 (0.156 - 79.778)	0.428
CRP	4.58 $\pm$ 3.89	19.3 $\pm$ 29.01	1.153 (0.928 - 1.432)	0.200	1.342 (0.703 - 2.561)	0.373	1.222 (0.77 - 1.939)	0.395
AST	53.4 $\pm$ 46.72	54.55 $\pm$ 57.53	1 (0.982 - 1.019)	0.966	1 (0.869 - 1.151)	0.995	1.003 (0.869 - 1.157)	0.970
ALT	24.6 $\pm$ 20.67	41.41 $\pm$ 73.03	1.008 (0.976 - 1.041)	0.642	1.003 (0.831 - 1.211)	0.975	1.017 (0.811 - 1.274)	0.886
Urea	17.6 $\pm$ 9.81	21.38 $\pm$ 16.29	1.029 (0.914 - 1.159)	0.635	1.34 (0.764 - 2.351)	0.307	1.282 (0.808 - 2.035)	0.292
Krea	0.4 $\pm$ 0.42	0.29 $\pm$ 0.2	0.216 (0.007 - 6.39)	0.375	0 (0 - 87.539)	0.171	0 (0 - 58.187)	0.160
pH	7.3 $\pm$ 0.12	7.33 $\pm$ 0.09	27.961 (0.001 - 687284.597)	0.518	---	---	---	---
pCO ₂	40.6 $\pm$ 7.13	42.29 $\pm$ 9.85	1.022 (0.911 - 1.146)	0.711	---	---	---	---
HCO ₃	20.54 $\pm$ 5.12	21.67 $\pm$ 3.69	1.07 (0.851 - 1.346)	0.561	---	---	---	---
BE	-5.54 $\pm$ 6.96	-2.88 $\pm$ 5.27	1.079 (0.921 - 1.265)	0.347	---	---	---	---
Lactate	3.92 $\pm$ 4.4	2.14 $\pm$ 0.98	0.712 (0.429 - 1.181)	0.189	---	---	---	---

¹Coc&Snell R²: %40,3; ¹Nagelkerke R²: %64,5; ²Coc&Snell R²: %39,3; ²Nagelkerke R²: %63

Alternative Treatment Methods and Application to Cardiovascular Areas with Summary of Some Related Studies

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Introduction

Heart attack is a heart disease with the highest morbidity and mortality rate among cardiovascular diseases and is very common in today's societies. The heart is a powerful organ that pumps blood an average of 60-80 times per minute during rest. It needs to be able to work on its own with the whole body and to feed itself for the continuity of this work. These self-feeding vessels are called coronary arteries (İlhan, 2016). If a deterioration occurs in these vessels, what we call ischemia occurs and the heart muscle, namely myocardium, cannot be fed adequately. And in this case, what we call myocardial infarction, that is, a heart attack occurs. Coronary insufficiency tables vary according to the type, degree and location of the stenosis in the coronary vessels. In some patients, there may be chest pain that occurs only during physical activity and is relieved by rest, as well as a heart attack (infarction) that sometimes develops as a result of sudden occlusion of the vessels and can lead to sudden death, starting with severe chest pain (Aybek, 2020).

When a part of the heart muscle undergoes ischemia and is deprived of oxygen, the myocardium cannot perform its function and a heart attack occurs. Patients' approach to alternative treatments has increased due to factors such as the expensiveness of the medical and surgical treatment used, the risky, the difficult accessibility for low and middle-income patients, and the greater confidence of people in natural methods. Although there are different methods in the treatment of heart attack in the world and in our country, research on alternative treatment has increased in many countries such as China and India, along with classical medical and surgical treatment. Among the alternative treatments, especially in many countries, especially in China, garlic, green tea, temple tree, licorice, cloves, ginger, St. John's Wort, pomegranate juice and cranberry juice are used as blood thinners, cholesterol-lowering, anti-hypertension, anti-atherosclerosis in the treatment of heart attack.

A heart attack usually has typical symptoms that manifest itself in the form of a pressing and compressive pain in the middle of the chest, radiating from the chest to the neck, spreading to the chin ears and arms, and especially numbness in the left arm and two little fingers (Şentürk, 2008). In humans, many factors such as age, obesity, genetic predisposition, alcohol and cigarette use, malnutrition, etc., form the basis for heart attack and many similar cardiovascular diseases and trigger disease formation.

However, among the important risk factors of heart attack can be counted as; Vascular diseases or heart diseases such as atherosclerosis, a previous heart attack, any rhythm disorder or faint, age (over 40 in men, over 50 in women), smoking, excessive alcohol consumption, some illegal drugs and derivatives; high levels of triglycerides, high bad cholesterol and low good cholesterol content, diabetes, high blood pressure, and chronic high stres (Jensen,1991). The diagnosis of heart attack is made by the symptoms seen in the patient, the patient's history, physical examination, EKG and blood tests (Şentürk, 2008).

Every year, approximately 18 million people die due to cardiovascular diseases and other diseases it causes. When the causes of death worldwide are examined, it has been determined that cardiovascular diseases are the number one cause of death in 2019 (Roth et al., 2020).

It is seen that most of these death rates are in middle-income countries such as Central Asian and Eastern European countries and some low-income countries. Heart attack is one of the diseases with the highest mortality and morbidity rates (Kılınç et al., 2021). According to the worldwide research statistics of the World Health Organization, the number of deaths from cardiovascular diseases is expected to reach 23 million in 2030 (https://nccih.nih.gov/research/statistics/2007/camsurvey_fs1.htm). According to the same report, 18 million people die every year due to heart diseases. According to the report of the American Heart Association, an increase of approximately 46% in heart failure is expected from 2012 to 2030; In 2030, it is predicted that more than 8 million people aged 18 and over in the USA (with 5.8 million in 2011) will have heart failure (Beştepe et al., 2020). It is understood that this estimated increase has triggered further research on heart attacks. Losses due to cardiovascular diseases in Turkey are equivalent to 30% of all deaths, and this rate represents half of the deaths from cancer (Coşkun et al., 2009). 31% of deaths in the world are due to cardiovascular diseases, and 85% of them are caused by heart attack and stroke. When we look at the 2018 Turkey data, 36.7% of the deaths are caused by cardiovascular system diseases, 39.1% of them are due to ischemic heart disease and 22.2% are due to cerebrovascular diseases (T.R. Ministry of Health, 2019).

Considering all these data and evaluations, it is seen that coronary heart disease and cerebrovascular disease are the most common causes of death among cardiovascular system diseases.

In addition, even though the deaths due to cardiovascular system diseases have decreased over the years, it is still seen that it is the most important cause of death both in the whole world and in Turkey, and the complications related to these diseases are increasing (Türk et al., 2021).

Considering the deaths caused by cardiovascular system disease, 36% of them are premature deaths occurring before the age of 75. In terms of chronic heart disease, cerebrovascular disease and peripheral arterial disease, atherosclerosis was found to be responsible for 71% of all deaths in 1997. When looking at the 35-45 age range in men, the incidence rate is 7% per year, while this rate rises to 68% at the age of 85-95. In women, the difference decreases as the age progresses, but if you look at the 10 years later, it is seen that the frequency increases at similar rates (İşlegen, 2007).

Many factors such as the increase in education level, increase in income levels, change in dietary habits and controlling infectious diseases in our country and throughout the world have led to an increase in life expectancy. The prolongation of the average life expectancy has significantly increased the percentage of individuals aged 65 and over in the general population in both developed and developing countries, and the calculations show that 20% of the world population will be 65 years and over in 2050 (Genç and Yiğitbaş, 2021). Among the alternative treatments, such as garlic, green tea, temple tree, licorice root, clove, ginger, St. John's Wort, pomegranate juice and cranberry juice, which are mainly preferred in many countries, especially in China, in the treatment of heart attacks, blood thinners, cholesterol lowering, antihypertensive herbal treatment sources used as a preventative against atherosclerosis. Despite all the developing opportunities, most of the patients die before they can even reach the health institution. Therefore, people should be informed about heart diseases, heart attack and its symptoms and should be directed to apply to the nearest health institution as soon as the symptoms are felt. Patients with heart attack symptoms apply to the emergency room, and treatment begins quickly. Oxygen support is provided by measuring the person's blood oxygen value. People with high pain can be benefited from various painkillers (Daşkapan, 2013). The main purpose of treatment; It is the removal and opening of the

blockage in the vessels feeding the heart and the immediate restoration of blood flow to that area. Tissue oxygenation is restored and myocardial function is restored. For this purpose, aspirin is given to remove blood clotting (blood thinner-anticoagulant therapy), nitroglycerin and oxygen therapy to relieve chest pain and improve blood flow.

CAM (complementary and alternative treatments)

In addition to all these medical treatments, as mentioned before, the tendency to alternative treatments has increased rapidly due to factors such as easier economic opportunities, accessibility for patients and their relatives, and the fact that natural methods create a more reliable impression on people. With this; Complementary and alternative medicine terms are generally gathered under one title and evaluated as a whole. Alternative medicine is defined as any health service that replaces medical treatments and is not included in modern biomedicine or treatments (Dömbekci et al., 2020). Complementary medicine, on the other hand, is defined as the treatment and care systems applied in addition to medical treatment (Ceyhan and Yiğit, 2016). According to the definition made by the United Nations National Institute of Health, “Complementary and alternative medicine; It is a broad field of health that encompasses all health services, methods, practices and accompanying theories and beliefs outside the politically dominant health system in a particular society or culture at a given time (Karayağız and Öztürk, 2008).

In other words, many definitions can be made about concepts such as complementary therapy, which are shaped by the culture, value judgments and beliefs of the society we live in. Traditional Medicine, or in other words, folk medicine, aims to evaluate the information to be obtained from the religious beliefs, socio-cultural values and other elements of the culture of the society in which it is transmitted from generation to generation and to use it in the treatment of diseases (Sütçü, 2018; Ersoy, 2014). An holistic approach to the concepts of health and disease is the essence of complementary alternative therapies. Observation, trial and error, and the use of datas obtained in this way as treatment have been transferred from each generation to the next and not only contributed to medical science, but also used as alternative treatment (Ersoy, 2014). The basis of alternative and complementary therapy applications is based on Chinese and Ayurvedic medicine treatments, and it has developed in many areas towards the end of the 19th century. The United States established alternative medicine offices within the national health institutes in 1993 and expanded its powers in 1998. It changed its name to the national center for alternative and complementary medicine (NCCAM, 2018). The purpose of this center; It is stated as to examine the reliability and effectiveness of complementary and alternative medicine applications and to ensure the participation of scientifically proven practices in traditional treatments (Karayağız and Öztürk, 2008).

However, when we summarize the factors that direct people to alternative treatments, the following come to forward;

- *effective and high reliability
- *The patient has a feeling that he can control the treatment
- *does not include invasive methods
- * therapists spend more time with the patient
- *besides being accepted easily and getting depressed
- * fear of side effects of applied medical treatments
- *insecurity
- *The patient has no hope of recovery.

(Ceyhan and Yiğit, 2016)

In general, the most common and most preferred complementary and alternative medical treatments from the past to the present; It covers ancient approaches such as Ayurveda, Acupuncture, Traditional Anatolian Folk Medicine, Herbalism and Yoga, and new approaches such as Oxygen Therapy, Ozone Therapy, Detoxification Therapy, Aromatherapy, Homeopathy, Dietary Support, Electromagnetic Fields, Massage, Reiki, Bioenergy and Hypnosis (<http://tr.wikipedia.org/wiki/Alternatif>).

Ayurveda treatment, which is the most preferred and widely used in countries such as China and India, is one of the oldest treatment methods (X century BC). It literally means 'science of wellness' and originates from the mystical philosophy of the Hindu religion.

It is widely practiced in Nepal, Sri Lanka and Bangladesh. It is accepted as focusing on healthy life instead of preventing the emergence of the disease or instead of intervening when the symptoms of the disease occur or curing the diseases. According to this system, the human body and the universe in which it lives are a whole, and human health is actually considered as a balance of various elements that make up this whole. Ways to live long and healthy; Recommendations on nutrition-lifestyle and physical-psychological health are presented in the form of complex systems such as massage, meditation and yoga techniques. Ayurveda practice can be summarized as maintaining a long and healthy life by reducing the effects of time and strengthening the body's immune system (Ozturk, 1990).

Another treatment method that attracts the most attention and has been the subject of research in recent years is herbal treatments. Since ancient times, plants have been used as medicine and serious successes have been achieved in the treatment of many diseases. Especially since the end of the 20th century, the tendency to treatments with herbs has increased rapidly due to reasons such as being more economical, easily accessible, and having less side effects and toxic effects. Herbal treatment sources such as garlic, green tea, ginger, Echinacea, Gingko biloba, Ginseng, Elderberry, Turmeric, St. John's Wort, Valerian, Chamomile are traditionally used to treat numerous illnesses, including heart disease, dementia, sleeping problems and some others. Because herbal remedies are derived from natural sources, they are generally considered safe, but this may not always be the case. Like traditional medicines, the use of herbal supplements may cause serious side effects or interact with other medicines taken.

When looking at complementary therapies in general, they are classified according to different criteria as follows:

1-Traditional Alternative Medicine: This field includes more common and accepted forms of therapy such as acupuncture, Ayurveda, naturopathy, homeopative, Oriental practices (Natural therapy, Chinese or Oriental medicine). Acupuncture is a kind of therapeutic model based on traditional Chinese medicine (includes Chinese herbal medicine, moxibustion, cupping, Chinese massage, mind-body therapies such as Qigong and tai chi, and dietary therapy). These therapies have been practiced around the world for centuries.

2- Biologically-Based Therapies

The biologically-based therapies include aromatherapy, diet-based therapies, chelation therapy, folk medicine, iridology, megavitamin therapy, phytotherapy/herbal medicine and neural therapy.

3-Body Therapies: Touch has been used in medicine since the early days of medical care. Tactile healing is based on the idea that illness or injury in one part of the body can affect all parts of the body. With manual manipulation, the body can fully focus on healing at the site of injury or illness if other parts can be brought back to optimum health. Body techniques are often combined with mind techniques. Examples of body therapies as chiropractic and osteopathic medicine; acupressure, Alexander technique, Bowen technique, chiropractic manipulation, Feldenkrais

method, osteopathic manipulation, reflexology, Rolfing, Trager bodywork, and Tui na, Meditation; Biofeedback and Hypnosis, Massage; including, Swedish massage, sports massage, deep tissue massage, and trigger point massage, body movement therapies; biofeedback, bioresonance, cognitive-behavioral therapies, deep-breathing exercises, tai chi, yoga, and shiatsu can be given. According to one of study; the use of frequency of this therapies change from 2% to 57%, with deep breathing and meditation most preferred. In this study, it was indicated that, Cardiac patients used these for stress, emotional health, and general wellness.

4-Dietary and Herbal Approach: For centuries, human beings have moved from a simple diet of meats, fruits, vegetables and grains to a diet consisting of foods usually rich in fats and complex carbohydrates. Both, as well as nutritional surplus and deficiency, have become problems that lead to certain chronic diseases in today's society. Many dietary and herbal approaches are used to balance the body's nutritional health. Within diet and herbal approaches, Dietary supplements; herbal medicines; Nutrition / Diet applications are included.

5-External Energy Applications: Some people believe that external energies from objects or other sources directly affect a person's health. Examples of external source energy therapies as Electromagnetic treatments; healing touch, light therapy, magnetic therapy, millimeter wave therapy, sound energy therapy, Reiki and Qigong can be given. Reiki include therapeutic touch and healing touch, known as together "hand-mediated" therapies and its aim to reduce pain and anxiety and increase health via healing energy aspect.

-Forms of Sense-Based Treatment: Some people believe that the senses, namely touch, sight, hearing, smell and taste, can affect overall health. Examples of therapies involving the senses are Art, dance and music; Visualization and auxiliary images also can be added (<https://www.hopkinsmedicine.org/health/wellness-and-prevention/types-of-complementary-and-alternative-medicine>; Rabito and Kaye, 2013).

In addition to all these, various researches on cardiovascular, lung and blood diseases and their treatments were conducted by the National Heart, Lung and Blood Institute and the National Institute of Complementary and Alternative Medicine in 2001.

In the workshop held on the general use of CAM applications, there were positive opinions on the fact that individuals began to prefer CAM applications to a large extent, and for this reason, a joint decision was taken to initiate large-scale clinical studies in order to maintain and protect the health of the population. It was emphasized that more serious research should be done to understand the basic functioning of CAM applications, that time and resources could be wasted and even risky. It is thought that meaningful basic and clinical studies are needed to standardize the use of procedural CAM interventions and herbal products in treatment (Lin, 2001).

Standart Conventional Treatment and Drugs for Cardiovascular Treatment

After the definitive diagnosis of heart attack is made, it is necessary to start treatment quickly to unclog the coronary vessels in the cardiovascular system, and for this purpose, two treatment methods are applied;

These are;

1. Clot Dissolving Drugs (Thrombolytic Therapy)
2. Percutaneous coronary intervention (coronary angiography and angioplasty)

If we look at the interaction between drugs used in the treatment of cardiovascular diseases and CAM;

-Warfarin and its use for antithrombotic therapy:

Many herbal uses have interactions with warfarin. During the use of herbal therapy, it increases the risk of bleeding in patients receiving warfarin or conventional antiplatelet therapy, etc. It can make positive or negative changes in platelet function, such as In many cases, it is thought to increase the likelihood of bleeding; however, numerous cases of increased bleeding tendency have been reported in patients using herbal supplements with or without warfarin.

-Digoxin: It has been explained that some plants can act as digoxin-like active substances and therefore, plants that are likely to act in this way can increase the digoxin effects of plants. Chinese silk vine (*Periploca Sepium*) is a poisonous plant containing cardiac glycosides. *Uzarae radix* (uzara root) has been noted to have cardiac effects like digoxin when used in high doses. Serious side effects of herbal treatments may result from lack of standardization and inadequacy in packaging and preparation.

-Cyclosporine: This substance, which is used for immunosuppression after heart transplantation, is exposed to many drug-drug interactions. For example, interaction with Diltizem causes increased levels of cyclosporine. If diltizem is used in patients with heart transplantation and hypertension, the dose of cyclosporine should be reduced. St. John's Wort is used as herbal treatment for those with depression and mood disorders. Unexpected interactions with cyclosporine have been reported in various sources. It is stated that it causes a decrease in the level of cyclosporine and its rapid elimination from the body (Miller et al., 2004).

After all these definitions of CAM and Cardiovascular area, we summarized many different CAM applications that have been tried in different geographies and including different applications in this field, and aimed to reveal the advantages and disadvantages of CAM applications in the Cardiovascular field, as well as being used in the treatment of many different diseases.

Summarized of some Studies on Cardiovascular Fields:

Studies to determine the use of CAM (complementary alternative therapy) in the field of treatment of heart disorders are also carried out in different countries.

One of them; It was conducted based on a questionnaire analysis between April 2013 and April 2014 to determine the use of CAM in patients in the Cardiac Surgery unit of Angers University Hospital in France. It was understood that 58% of the 154 patients included in the study used a complementary and alternative medicine at least once in their lifetime, 38% in the previous year before the disease appeared, and 14% between consultation and surgical intervention.

In total, 71% preferred alternative treatment at any stage of their illness, and only 29% of those who used it informed their doctors and paramedics about it. It has been understood that the use of complementary and alternative medicines is common among patients who apply to cardiac surgery units, but the authorities are not sufficiently informed about this issue (Dalmayrac et al., 2016).

Cardiovascular diseases (CVD) have traditionally been treated with medications and lifestyle changes. The use of complementary therapies in this field has been increasing in recent years. The use and success of CAM in the prevention and treatment of CVD, including hypertension, hyperlipidemia, coronary artery disease, heart failure, and arrhythmias, was discussed in this another study. Especially It has been suggested that Garlic, which is included in TAT herbal applications, may exert an antiatherosclerotic effect by interfering with the inflammatory and oxidative pathways and inhibiting the lipid shield in the vessels, and it may mediate decreased peripheral vascular radiation, vasodilation and smooth muscle expansion.

It has been concluded that alternative methods such as special nutrition programs, meditation, yoga, acupuncture, herbal herbs and vitamin supplements that can be used as an adjunct to traditional medical care to improve cardiovascular health and quality of life can be used in this area

as promising, and it has been emphasized that similar results have been obtained from different studies. But It was stated that physicians interested in the subject should be informed about the possible effects of CAM interventions such as diet, exercise, supplements, herbs and mind/body techniques (Aggarwal et al., 2017).

In the surgically created heart attack model, animals were administered T3 and T4 hormones via gavage as a treatment model in another study and the values of proapoptotic proteins p53, JNK, BAX, Bcl2 were examined to determine the effect of the treatment given afterwards. It was determined that thyroid hormones improve cardiac function and decrease the expression of proapoptotic proteins after heart attack since apoptosis is determined by genetically p53 evaluation. It was concluded that hormone therapy, which is one of the CAM methods, has advantages in terms of the therapeutic effect of heart attacks and the availability of alternative treatments against heart attacks (Castro et al., 2016).

In the heart attack model caused by doxorubicin carried out by Shahid et al. in 2021; It has been reported that doxorubicin was given every other day and intraperitoneally for two weeks, and oral administration was applied for 14 days to determine the cardioprotective effect of boswellic acids for treatment purposes. It was stated that blood lipid profile and cardiac biomarkers including LDH, CK-MB, CPK, SGOT and troponin T were measured and histopathological evaluations were made. According to the results obtained, it was stated that there was a significant increase in cardiac enzyme levels and full lipid profile parameters in the diseased group compared to the control group. On the basis of pretreatment with boswellic acids reduced the level of all measured parameters and reduced the severity of myocardial damage, as supported by histopathological studies as well (Shahid et al., 2021).

In a study conducted by Saravanan and Prakash in 2004 to investigate the effect of garlic (*Allium sativum*) on lipid peroxidation in heart attack; Iso was used to create a heart attack, and garlic oil was used as a treatment agent within herbal treatment CAM. As stated in the study; In myocardial necrosis caused by isoproterenol; A significant increase in serum iron content, a significant decrease in plasma iron binding capacity, ceruloplasmin activity and glutathione (GSH) level were observed, while a significant increase in lipid peroxide levels was obtained. It has been stated that the activities of antioxidant enzymes such as superoxide dismutase (SOD), catalase (CAT), glutathione peroxidase (GPX), glutathione-S-transferase (GST) and glutathione reductase (GRD) are significantly reduced in the heart with isoproterenol-induced myocardial necrosis. According to the results obtained; It has been stated that garlic oil provides a significant return with its curative effect in metabolic changes due to heart attack caused by isoproterenol. It is an important study in terms of using garlic-containing oil as a treatment agent and demonstrating the curative effect of garlic as an alternative supportive treatment (Saravanan and Prakash, 2004).

In the study of Mnafigui et al. in 2016; In the iso-induced heart attack model, the antithrombotic, anti-inflammatory and cardiac preventive effects of eugenol were investigated. There were first negative and then positive changes in troponin-T, CK-MB and LDH and ALT plasma levels in all disease-induced and treated groups. An improvement was observed in the markers indicating cardiac damage in the eugonal treatment group, and that eugenol reduced inflammation by reducing inflammatory mediator proteins; indicating eugenol has a high level of protective effects in the cardiac formation process. It is important that the use of eujonal as a treatment agent and the subsequent improvements in biomarkers reveal the curative effect of eugenol, the active ingredient of garlic, as a herbal CAM application (Mnafigui et al., 2016).

In the study conducted by Ganguly and Alam in 2015 to determine homocysteine and its effects on the cardiovascular system; It has been stated that homocysteine can initiate an inflammatory response in vascular smooth muscle cells by stimulating the production of CRP. These findings provided new evidence for the role of homocysteine in the pathogenesis of the

disease and it was stated that it may be sufficient to prove the role of homocysteine in the pathogenesis of cardiovascular disease.

In this study, it was found that hyperhomocysteinemia may develop due to folate, vitamin B12 or choline deficiency, and It has been stated that the negative situation can be corrected and the risk of CVD caused by it can be prevented with supportive treatments such as nutrition and exercise. This study is important in terms of revealing the effect of homocysteine use and exposure to hyperhomocysteinemia on cardiovascular disease risk factors such as myocardial damage and also the effect of folate, vitamin B12 or choline deficiency on the occurrence of heart attack, which can be complemented by CAM supplementation applications. (Ganguly & Alam, 2015).

In other study; 96 male Wistar rats (16 weeks old) were used by dividing into 12 groups. Transthoracic echocardiography was performed on the rats. 4 weeks' treatment was applied with using *Allium sativum*, *Peganum harmala* and *Apium graveolens* as CAM methods. According to findings, *Allium sativum*, *Peganum harmala* and *Berberis vulgaris* could significantly improve cardiac function by improvement of left ventricular remodeling, lowering hs-CRP and NT-ProBNP and echocardiographic indexes without toxic effect into rats, suggesting they can be used as an alternative treatment with no harmful effect (Keihanian et al., 2021).

Oxidative stress is considered an important factor for several different disease including cardiovascular disease (CVD). An active components of Chinese herbal medicines (CHMs) have been searched and widely used for the treatment of CVD, including hypertension and coronary heart disease. In a different study, various active CHM components have been searched and It has been found that the active CHM components including astragaloside, ginsenoside and resveratrol can regulate and modulate the oxidative stress and the circulatory system, suggesting they can be used as an potential drugs for the treatment of various CVDs (Chang et al., 2020).

Gallic acid is a trihydroxybenzoic acid found in tea leaves and some plants. In a other study, the effect of gallic acid as an herbal treatment, on cardiac dysfunction and fibrosis in a mouse model of pressure overload-induced heart failure and in primary rat cardiac fibroblasts were searched. It was found that Gallic acid reduces cardiac hypertrophy, dysfunction, fibrosis and the expression of collagen type I and connective tissue growth factor; It decreases left ventricular end-diastolic and end-systolic diameter, perivascular fibrosis, and recovers the reduced fractional shortening and it suppresses the expression of atrial natriuretic peptide, brain natriuretic peptide, skeletal α -actin, and β -myosin heavy chain. According to these findings; gallic acid is a therapeutic agent and can be used safely for cardiac dysfunction and fibrosis in chronic heart failure (Jin et al., 2018).

In a different study, heart attack induced by experimentally giving the active substance (isopretrenol) to rats and to cause myocardial damage by adding homocysteine to the drinking water of the rats. Resveratrol, garlic extract and euganol as a complementary treatment were given. Regarding heart attack and damage, experimentally administered isopretrenol causes heart attack and homocysteine causes myocardial damage, the applied active substances of CAM methods had a protective and therapeutic effect at different doses and levels, suppresses apoptosis on the basis of genetic, molecular and biochemical analysis, indicating they can be used effectively and safely (Kocaman, 2022; Kocaman and Dosay-Akbulut, 2021).

In different study, a cross-sectional survey was conducted to the patients of cardiovascular clinics at Singapore for 2 months in 2014. Information included subjects related to various aspects of CAM use and its effect. According to findings; The prevalence of CAM use was 43.4 % (333/768). Biologically-based systems (29.4 %) was the most preferred type of CAM. Some patients (19.0 %) used multiple types of CAM. Approximately half of patients (50.8 %, n = 169) found CAM effective in treating their heart condition, while a smaller proportion (25.2 %, n = 84) found CAM more effective than conventional western medicine in treating their heart condition. A small

number of CAM users (7.8 %, n = 26) come a cross with side effects from CAM which included gastrointestinal complaints, allergy and rash. These findings suggested that CAM methods are popular within and can be prefer and use safely by cardiovascular patients (Teo et al., 2016).

Another similar study was conducted to evaluate the effects of Mind-body practices (MBI) (eg, Tai Chi, yoga, meditation) included in CAM on individuals with heart failure. For this purpose, the questions were asked to 1314 participants for determining the use and effects of 9 CAM methods covering mind-body practice. The results were found as follows; Tai Chi (n = 7), yoga (n = 4), relaxation (n = 4), meditation (n = 2), acupuncture (n = 2), biofeedback (n = 2), stress management (n = 1), Pilates (n = 1) and reflexology (n = 1). Most of the participants (n = 22, 95.8%) experienced quality of life (14/14), exercise capacity (8/9), depression (5/5), anxiety and fatigue (4/4), blood pressure (3/5), stated that there was an improvement in criteria ranging from low to moderate according to heart rate (5/6) and heart rate variability (7/9) criteria. According to these data; It has been stated that CAM applications can create moderate advantages in people with heart disease and can be applied and preferred as a supportive treatment. (Dosay-Akbulut, 2020).

In a similar different study, 10,572 respondents with cardiovascular disease were used and analyzed data on their CAM use. Among them, 36% had used CAM in a 12 months period. The most commonly used therapies were herbal products (18%) and mind-body therapies (17%). Among herbs, garlic, echinacea, ginseng, ginkgo biloba, and glucosamine with or without chondroitin were preferred mostly. Among mind-body therapies, deep-breathing exercises and meditation come forward. The patient that used CAM for their cardiovascular condition within this survey, reported the therapies as beneficial (80% for herbs, 94% for mind-body therapies) (Yeh et al., 2006).

Another study; based on determination of CAM using ratio after acute myocardial infarction (AMI). Patients with AMI were assessed using Angina Frequency and Quality of Life domains from the Seattle Angina Questionnaire and the Short Form-12 Physical and Mental Component scales for their CAM using ratio and their choosing categories (as mind-body, biological, and manipulative therapies) prior to and 1 year after AMI obtained from 24 US sites.

Among 1884 patients which they indicated not using CAM at the time of their AMI, 33% of them reported that started at least 1 forms of CAM therapy 1 year following AMI. 62% adopted mind-body therapies, 42% adopted biological therapies, and 15% started to use manipulative therapies. According to findings; they were not obtaine any association between CAM use and health-status recovery after AMI within patients of this study (Shafiq et al., 20016).

Coronary heart disease is determined and known with the formation of arterial plaque. If care is not taken, it can be cause important problems like myocardial infarction and heart failure. Zhishi Xiebai Guizhi Decoction can be use as a representative prescription in the treatment of coronary heart disease. This another study was dizayn to give the information related to mechanism of Zhishi Xiebai Guizhi Decoction in treatment of coronary heart disease. In this study, first of all the content of Zhishi Xiebai Guizhi Decoction was determined. Then, it was aimed to determine and analyze the molecular targets of Zhishi Xiebai Guizhi Decoction in coronary heart disease by using OMIM and GeneCards databases.

In the findings; by using network pharmacology analysis, they were found that different important core compounds of Zhishi Xiebai Guizhi Decoction (includes five herbs; as *Aurantifructus immaturus*, *Allii macrostemonis bulbus*, *Trichosanthisfructus*, *Magnoliae officinalis cortex*, and *Cinnamomi ramulus*and) has therapeutic efficacy and anti-inflammatory effects especially agains to chest stuffiness and indirectly against to cardiovascular diseases.

According to results; they indicated that Zhishi Xiebai Guizhi Decoction as one of CAM types may play a role in treating coronary heart disease with successfully via several different signaling

pathways and it can be nominated in using against to coronary heart disease (Liu et al., 2022; Gao et al., 2021).

Another study; carried out to determine the using ratio of CAM within hospitalized patients with acute coronary syndrome from March 1, 2001 to October 31, 2002. Because the complementary and alternative medicine (CAM) is generally preferred by patients with cardiovascular disease. Poisson regression models were used to assess the results.

In the findings; CAM using ratio was 19% in patients with coronary artery disease (CAD). Most of them who used CAM were non-Caucasian (31% vs 12%), uninsured (12% vs 7%), economically burdened (58% vs 29%), and had depression (13% vs 6%, $p < 0.05$ for all). The most preferred CAM types in patients using CAM are beta-blockers (64% vs. 46%, $p = 0.008$), aspirin (73% vs. 74%, $p = 0.90$), and statins (68% vs. versus 71%, $p =$).

According the results, CAM users of this study have worse socioeconomic status but in comparison with standart therapies no evidence was obtained that using of CAM cause to worse situation (Decker et al., 2007).

There is unsufficient datas on the use of complementary and alternative medicine (CAM) by patients diagnosed with coronary heart disease (CHD). In this study; the using ratio of CAM among patients with CHD, the reasons and factors effecting their choise, the preffered types of CAM, and the relationship between CAM using situation and patient's demographics were searched.

A cross-sectional questionnaire descriptive study was carried out with Palestinian CHD patients from three different hospitals. Siahpush scale was used to determine the approach of CHD patients toward CAM use.

150 patients were interviewed and 128 (85.3%) of the them completed the questionnaire. The majority of CAM users indicated they use CAM for health problems other than CHD, while a total of 59 (45.9%) patients have used CAM for their heart problems. Also; religious practices were found to be the most preferable form of CAM used by patients, while body and traditional alternative methods were the less.

According to the findings; Participants of this study commonly prefer CAM. Also, Religious practices come to first (Salah et al., 2020).

Another study was carried out to determine the possible use of Traditional Chinese medicine (TCM) within the first 24 hours of hospitalization (early IV TCM) for acute myocardial infarction (AMI). Datas were collected from 175 modern Western medicine hospitals throughout China including total of 14 097 patients with AMI across the 3 years. Within the CAM techniques the most preferable ones was herbal medicines and divided into 6 main groups according to their contents. These are respectively, *Salvia miltiorrhiza*; *Panax notoginseng*; *Folium ginkgo*; *puerarin*; and *breviscapinun*. According to result, almost all (99%) hospitals tried early intravenous TCM. The most commonly used early intravenous TCMs were *Salvia miltiorrhiza* (danshen) (35.5%), after that *Panax notoginseng* (12.2%); *Folium ginkgo* (7.5%); *puerarin* (2.7%); and *breviscapinun* (2.2%) come to forward respectively. On the basis of analyses, no significant associations between early intravenous TCM and in-hospital bleeding or mortality was obtained (Spatzs et al., 2018).

Complementary and alternative medicine (CAM) is most commonly used in China in the treatment of coronary artery disease (CAD). In different study; to determine the CAM using ratio and types and related informations against to CAD, a questionare study was design in Beijing. The obtained datas were analyzed with Student's t-test and chi-square test. The datas wete obtained from 546 patients with a diagnosis of CAD in the period of May to July, 2009. According to findings; 69.1% of the patients with CAD used at least one of CAM types. Patent herbal medicine (90.7%) was the most preferred CAM type. Comparison to non-CAM users, It was reported that

CAM users were older ($p < 0.01$), had a longer disease course ($p = 0.02$), and had better current health status.

On the basis of results; there are several different and important factors which can affect the use of CAM by CAD patients (Chu et al., 20013).

In another study; the use of complementary and alternative medicine (CAM) possibilities, ratios, types and the reasons among people with type 2 diabetes and/or cardiovascular disease (CVD), and the effect onto the quality of life were searched.

For this study; a survey was applied via mail and internet in Victoria, Australia.

The applicants were adults with type 2 diabetes and/or CVD, questiones were asked about whether they used CAM therapy as well as conventional medical treatment.

In the findings; The answers were given as follows; from 2766 people, 45.1% had used CAM in last 1 year period with between 20-96 years old ages. CAM users reported lower (worse) quality of life data than nonusers. In contrast, CAM users reported more positive changes in their perception of illness and behavior towards it than nonusers.

Most CAM users (94.6%; $n=1180$) used CAM, primarily vitamins, minerals and other nutritional supplements; 96.9% used them in addition to prescription drugs in standard treatment. 54.5% ($n=671$) of CAM users received counseling from CAM practitioners.

The rate of recommendation for the first use of CAM by a doctor, pharmacist or another healthcare professional (42.7%); Belief that CAM will be good (39.4%); The belief that CAM is not harmful (33.3%); a desire to have more control over their health (31.3%); desire to use “natural” (27.8%); close circle advice (23.8%); results were obtained as dissatisfaction with the standard treatment (18.2%).

These findings show that the use of CAM has a positive effect on the perspectives of people with chronic conditions. Considering the general high use of CAM, there seems to be a belief that the use of CAM can support the prevention of chronic diseases (Canavay et al., 2013).

Complementary and alternative medical (CAM) therapies are growing in popularity. There is little information on the use and success of CAM therapy, especially in patients with cardiovascular disease (CVD). In another study; telephone questionnaires were applied to 107 randomly selected patients with CVD. In the findings; 64% of the surveyed patients reported using CAM. Nutritional supplements (40%) and vitamins (35%) were the most preferred ones. Most of the CAM users (65%) declared that they applied to CAM because of their heart conditions. Friends or relatives (43%) or their own doctors were reported as the sources of information about CAM. However, the majority of respondents stated that they also received an oral supplement at the same time as CAM.

According to findings; It is seen that the use of CAM treatments is high on the basis of patients participating in the survey. Therefore, more illumination and studies are needed on this subject (Wood et al., 2003).

The use of complementary and alternative medicine (CAM) is increasing day by day. People with cardiovascular diseases frequently use CAM, despite significant side effects and insufficient information to support its use. This study was planned to reveal the relationship between CAM use in heart patients, the types of CAM used, the reasons and factors affecting its use, patient demographics and CAM use.

This study was conducted with 329 adult heart patients in the South-West District of Trinidad and Tobago. A questionnaire was applied to the participants between July 1, 2012 and August 31, 2012. Data analysis included χ^2 tests and binary logistic regression.

In the result; One hundred and eighty-five (56.2%; standard error [SE] = 2.74%) patients were found to use CAM. Herbal medicines are the most common (85.9%; SE = 2.56%); Among the CAM methods used, spiritual therapy/mind-body systems (61.6%; SE = 3.58%), physical therapy/body manipulation (13.5%; SE = 2.51%), alternative systems (8%, 1; SE = 2.01%) and other methods (3.8%; SE = 1.41%) respectively.

The most frequently referenced CAM subtypes were determined as omega-3 fatty acids (54.6%), B vitamins (50.4%), special diet/nutritional supplements (41.1%), and faith healing (54.1%). Participants believe that CAM improves health and wellness (79.5%; SE = 2.97%), is supportive in the fight against diseases (78.9%), believes that it can overcome the limitations of standard treatment (69%), those who believed it relieved symptoms (21.6%), cost less than standard treatment (CM) (21.6%) and had symptoms those who stated that they improved less than those who stated that there were fewer adverse/damaging effects compared to standard treatment (CM) (29.7%) or those who were unsatisfied with CM treatment (12.4%) results are declared. Ethnicity has been found to have an impressively greater effect on CAM use than religion.

According to findings; In this study, the use of complementary and alternative medicine was found to be high in heart patients (56.2%, SE = 2.74%), and it was revealed that ethnicity and close environment affected the use of CAM (Bahall, 2015).

It would be beneficial for patients with coronary heart disease to learn how to manage their condition and make the necessary controls in order to maximize their quality of life and prevent recurrence or worsening of the disease. This study was planned to investigate the behaviors of patients towards the use of complementary and alternative drugs and therapies (CAM) within 1 year after experiencing a cardiac problem in England.

A questionnaire was applied to 463 patients who applied to 4 hospitals in the West Midland region of England during a 12-month follow-up period. In the results; 91.1% of the patients participated in the survey study and 29.1% stated that they used CAM (mostly, acupuncture, homeopathy, chiropractic, dietary supplements, massage and rubbing for aches and pains, exercises) and/or one of the test kits, but very few (8.9%) used both methods. According to the results, CAM was used more frequently in the treatment of other diseases than cardiovascular diseases. Self-test kit use (77.2%) was found to be more common than CAM (31.7%), the most common (80.0%) being recorded as blood pressure monitors. The majority of the patients (89.5%) declared that they bought these test kits from various sources on their own initiative.

On the basis of findings; It was stated that the patients tried new technologies to monitor their cardiovascular health, and the rate of CAM usage was less than that of test kits (Greenfield et al., 2008).

This another study was conducted to evaluate the frequency and types of CAM treatments in patients hospitalized with acute coronary syndrome arising from cardiac causes.

An attempt was made to reveal the profile of CAM users. Data were collected through interview and based on information on demographics, patient's medical history, CAM use, doctor visits, and patient-physician communication. 223 patients provided the necessary information by meeting the appropriate criteria and filling out the questionnaire. Sixty-three percent stated that they had used at least one CAM treatment in the past 1 year for general health purposes. Only 11.7% of the patients reported using CAM for cardiovascular problems. The rate of CAM use was found to be higher in women than in men, and it was observed that they tended to use more CAM treatments. More than one third (35.9%) of the patients using CAM did not inform their doctor about it (Barraco et al., 2005).

Complementary and alternative medicine (CAM) can provide both risks and benefits for people with cardiovascular disease. In this study,

A general systematic study was conducted, including studies on the use of CAM in patients with cardiovascular disease. Twelve of the 27 studies included in this overall evaluation were in the United States; six of them were made in Canada. The remaining nine were conducted in England, Hong Kong, India, Italy, Korea, Nigeria, Spain and Turkey. It was stated that sample sizes ranged from 65 to 10572 patient data.

When we look at the results; Twenty-seven studies were included in this study and the data were evaluated. It has been observed that the rate of CAM use in heart patients varies between 4-61%. Among the CAM types used, the rate of biologically based treatments varied between 22% and 68%, while herbal medicines were used between 2% and 46%. Within the biologically based treatments; vitamins, minerals and other dietary supplements were preferred and used between 3% and 54%. The most used supplements were vitamin B/B12 or B complex, vitamin C, vitamin E, calcium, glucosamine/chondroitin, coenzyme Q10, calcium and magnesium. The most preferred herbs were echinacea (2% to 14%), garlic (1% to 69%), ginger (1% to 24%), ginkgo (1% to 9%), and ginseng. The using ratio of mind-body therapy (MBT) ranged from 2 to 57% with prayer and fasting. Aiming good health were the main reasons for taking vitamins and herbal therapy as well as specifically to treat cardiovascular disease and believe to get benefit from CAM. The majority of patients did not inform their healthcare professionals about their use of CAM.

According to these results; The use of CAM appeared to be common in patients with cardiovascular disease. The findings stated that the effects of CAM on the medical treatment of cardiovascular disease were not taken seriously and patient-physician communication should be strengthened in this regard (Grant et al., 2012).

Another study was conducted to examine the frequency and pattern of complementary and alternative medicine (CAM) use in patients with hypertension who applied to a hypertension clinic. 225 patients who applied to the hypertension clinic of Lagos University Teaching Hospital over a 3-month period were interviewed and data were collected. Information on socio-demographic data, hypertension, blood pressure, compliance with conventional drugs and use of CAM were obtained. Healing systems, mind-body connections, dietary supplements and herbs, energy therapy, and manipulation and touch therapy types were classified into questions to determine the CAM methods used.

When we look at the results; Data from 90 (40%) male and 135 (60%) female patients were evaluated. 88 (39.1%) of the participants stated that they use CAM. Herbal products were declared as the most preferred type of CAM. Garlic was preferred as the most frequently used herbal product (69.3%). Others are native plants (25%), ginger (23.9%), bitter leaf (*Vernonia amygdalina*) (9.1%) and aloe vera (4.5%). Spiritual therapy was used at a rate of 2.5%. None of the patients used healing systems, energy therapy, or manipulation and touch as CAM types. There was no difference between CAM users and non-users in terms of characteristics such as socio-economic status and blood pressure control.

According to these results, it has been revealed that a significant proportion of hypertensive patients use CAM therapies. Due to the high rate of use of herbal CAM methods among the hypertensive patients participating in this study, it is seen that it is important to inform the physicians about the benefits and harms of such practices for hypertension and related heart diseases that have the potential to occur (Amira et al., 2007).

In another similar study; Information was obtained from 200 randomly selected adults by face-to-face interviews in Durban, South Africa. When we look at the results;

The prevalence of CAM use for the 2000/2001 period was found to be 38.5%. Spiritual healing and herbal/natural medicines, including vitamins, were the most common types, 42.8% and 48.1%, respectively, within the general CAM use. People stated that they use CAM for the treatment of

diseases such as diabetes mellitus, headaches, arthritis and joint pains, stress, skin disorders, backaches, hypertension and nasal disorders, especially cardiovascular disorders. Half of CAM users also preferred drugs. Analysis showed that age, gender, marital status, religion, education level and income did not affect the use of CAM. More than half of CAM users (51.9%) stated that they used it after a relative recommendation or a CAM-related advertisement in the press. Seventy-nine percent of users stated that they got positive results from their treatments. Fifty-four percent of CAM users (excluding those who only use spiritual healing) said that they did not inform their doctors about using CAM.

Although CAM is preferred to treat many different ailments, this practice is not specific to a particular demographic and has been found to be similar to the rest of the world in many ways. It was stated that since most CAM users are satisfied with the effects of CAM, it would be beneficial to conduct more research on this subject and reveal its possible positive and negative effects (Singh et al., 2004).

Another similar study was conducted to evaluate the current and prospective use of complementary and alternative medicine (CAM) therapies in outpatients with cardiovascular disease. This study was planned in response to the increasing interest in CAM treatments and the need to determine their use, especially by patients with cardiovascular disease. The data was created with a 17-question questionnaire. The questionnaire was completed by 1,055 patients (655 males, 351 females; mean age 63.5 years). Of the respondents, 36.8% had cardiac symptoms for more than 10 years, 48.2% had coronary artery disease, and 82.5% stated that they had used CAM therapy. In the use of CAM, as a variety; 75.4% stated that they use dietary supplements, 31.5% chiropractic treatment, 23.9% mind-body therapies and 19.2% massage. The top 5 dietary supplements used were multivitamins (52.8%), fish oil (45.2%), vitamin C (36.1%), vitamin E (29.0%), and fiber (27.1%). The top supplements used for heart-related conditions were fish oils, omega 3 fatty acids, multivitamins, folic acid, coenzyme Q10, and vitamin E. Only 14.4% said that they informed their doctors about the use of CAM. The most preferred practices for improving cardiac symptoms were relaxation techniques, stress management, meditation and guided imagery. Among patients with coronary artery disease, the greatest benefit from using these methods was reported to be chest pain (20.0%), sleep disturbance (17.9%), leg swelling (17.9%), and shortness of breath (13.9%). It was stated that the majority of the participating patients used CAM therapies and were warmly welcomed to these applications (Prasad et al., 2013).

Conclusion

One of the diseases with the highest mortality rate in the world is heart attack. The incidence of ischemic heart disease, hypertension, diabetes and other chronic diseases, increased consumption of foods high in fat and sugar, and sedentary behaviors cause an increase in cardiovascular problems. In order to correct this situation where many risk factors are effective, to increase the quality of life of the society, to reduce the morbidity and mortality rates, and most importantly, for people to access the treatment in the most effective, fastest, most economical and most accessible way, there are different methods for the treatment of Cardiovascular problems.

Due to reasons such as the increase in the cost of classical drug therapy and surgical interventions in the world, the difficulty of reaching the treatment on time and within the available possibilities, the increase in awareness about the increasing side effects of drug therapy, it is at the forefront in developed and developing countries and is preferred as an alternative to other treatments. The use of alternative treatments has become widespread.

It is known that there is a significant increase in the preference of alternative methods, especially in the treatment of heart attack, in addition to the known anticoagulant and antithrombotic treatments. It is stated that early diagnosis and treatment of a heart attack reduces the damage to the heart, and with this, before a definitive heart attack is diagnosed, various

treatments are applied even in case of doubt to reduce the negative effect. Because of the narrow therapeutic range of cardiac drugs, it has been stated that the possibility of adverse interactions with dietary supplements and herbal medicines is of concern, therefore it is important to understand the frequency and nature of CAM use in patients with CVD in order to promote beneficial CAM therapies and prevent possible herb-drug interactions.

In this study, myocardial infarction, within the Cardiovascular problems and its alternative treatments, the usability of these treatments, as well as the efficacy and causes of alternative treatments in the world with summarize of some related studies are mentioned. It was concluded that complementary alternative treatment methods can be used in cases where conventional treatments are insufficient and under the control of a doctor.

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Current Treatment Approach in Mushroom Poisoning

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Introduction

Mushrooms are living organisms that are widely found in nature, some of them are consumed as food. Consumption of some types of mushrooms can lead to poisoning, which can even lead to death. Humid environments provide mushrooms to thrive. For this reason, mushroom poisoning is mostly caused by the consumption of mushrooms collected in rainy periods and regions.

90-95% of deaths due to mushroom poisoning are caused by amatoxins (Barbato, 1993; Diaz, 2018). Therefore, in this study, poisoning due to amatoxins and their treatment will be explained.

Amatoxins are oligopeptides and are selective inhibitors of RNA polymerases. Amatoxins are not inactivated by boiling or freezing. Alpha amanitin is the most hepatotoxic among amatoxins, and its lethal dose in humans is 0.1 mg/kg (Baumann, Münter & Faulstich, 1993).

Amatoxins absorbed from the gastrointestinal tract by ingestion of poisonous mushrooms come to the liver, which is the main target organ, via the portal vein (Karlson-Stiber & Persson, 2003). 60% of the alpha amanitin absorbed from the gastrointestinal tract is excreted in the bile, and some of it comes back to the liver by enterohepatic circulation (Busi et al., 1979). Amatoxins are taken to the liver by active transport, OATP (Organic anion transporting polypeptide) 1B3 and NTCP (Sodium-taurocholate cotransporter) (Letschert et al., 2006). Since messenger RNA cannot be synthesized due to RNA polymerase inhibition, protein synthesis and metabolism at the cellular level in the liver cease (Karlson-Stiber & Persson, 2003; Kröncke et al., 1986).

Most deaths caused by ingestion of mushrooms containing amatoxin are due to hepatocellular necrosis (Diaz, 2005). In the kidneys, amatoxin is filtered from the glomeruli and reabsorbed from the proximal tubules, so it can develop acute tubular necrosis in the kidneys (Gibbons, 1982). The rate of binding of amatoxins to proteins is quite low. They are excreted in urine and feces, and can be detected by high-performance liquid chromatography (Jaeger et al., 1993).

Most edible mushrooms are not toxic or cause mild or moderate gastrointestinal toxic effects. (Trestrail, 1991). The clinical manifestations of amatoxin poisoning are divided into four stages. The first stage is the latent phase, which is usually asymptomatic, lasting 0 - 24 hours. The second stage is the gastroenteritis phase, which starts approximately 6 hours after the mushroom is taken, and symptoms such as nausea, vomiting, diarrhea, abdominal pain, and hematuria are observed (6-24 hours). The third stage is the relative recovery phase in which asymptomatic elevations of hepatic enzymes are observed (24-72 hours). The last stage (4th - 9th day) is the hepatic failure phase in which acute liver failure develops, renal damage, encephalopathy, coma and death can occur (Berger & Guss, 2005). Symptoms seen after a prolonged latent period are more suggestive of amatoxin poisoning (Diaz, 2005).

For the diagnosis of mushroom poisoning, identification of the eaten mushroom, the time elapsed between the mushroom ingestion and the onset of the specified symptoms, and laboratory tests are used (Zilker, 2008). In case of suspected amatoxin poisoning, first of all, examinations

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such as complete blood count, blood glucose, urea, creatinine, electrolytes, aspartate aminotransferase (AST), alanine aminotransferase (ALT), bilirubins, prothrombin time (PT), active partial thromboplastin time (aPTT) and complete urinalysis should be done. For diagnosis of amatoxin poisoning, alpha amanitin can also be checked in the urine (Butera et al., 2004).

Prevention from Mushroom Poisoning

Mushrooms that grow in nature are collected in many countries for consumption as food. For this reason, mushroom poisoning is common and important health problem, especially in rainy periods and in rural areas. People who do not have sufficient knowledge and experience to avoid mushroom poisoning should not collect mushrooms and should not consume the mushrooms they collect. People especially in rural areas should be educated on this issue.

Treatment

Symptomatic patients should be hospitalized, supportive treatments such as fluid and electrolyte support and symptomatic treatments should be given. In the stage of hepatic damage, glucose-containing fluids should be preferred because of the risk of hypoglycemia (Ward et al., 2013).

Gastric lavage should be performed in patients presenting at an early stage with suspected mushroom poisoning. After lavage, it is recommended to administer repeated doses of activated charcoal (initial dose 1 g/kg, maximum 50 g in adults and 0.5 g/kg repeated doses) (Diaz, 2005; Zellner et al., 2019). Maximum benefit is obtained if activated charcoal is started within the first 24 hours after mushroom ingestion. However, in the first 4 days, 0.5 grams/kg can be applied every 4 hours. The biliary excretion of amatoxins can take up to 5 days. Activated charcoal binds amatoxins and allows them to be excreted in the faeces (Jaeger et al., 1993). There was a significant decrease in mortality in those who received repeated doses of activated charcoal and supportive treatment compared to those who received only supportive treatment (mortality 10% and 47%, respectively) (Enjalbert et al., 2002).

The most frequently used drugs in treatment are n-acetylcysteine, intravenous benzathine benzylpenicillin (penicillin G benzathine), silymarin and cimetidine (Karlson-Stiber & Persson, 2003).

N-acetylcysteine is an antioxidant and free radical scavenger. However, it has been proven that it is not hepatoprotective in cases of acute liver failure other than paracetamol, such as mushroom poisoning (Karlson-Stiber & Persson, 2003; Sklar & Subramaniam, 2004). However, it can be used in treatment considering its low side effects and potential benefits. In one study, it was found that the combined use of silibinin and N-acetylcysteine increased survival (Bergis et al., 2012).

High-dose (300,000 - 1,000,000 units/kg/day) intravenous penicillin has been shown to inhibit the uptake of amatoxin into hepatocytes. Benzylpenicillin acts by inhibiting the carrier protein of alpha amanitin (OATP 1B3) in the hepatocyte membrane (Karlson-Stiber & Persson, 2003; Magdalan et al., 2009).

Silymarin is one of the most studied drugs for the treatment of poisoning by amatoxins. However, no randomized controlled trials have been conducted (Schneider, Borochovit & Krenzelok, 1987). Silymarin is an antioxidant and free radical scavenger derived from thistle, which maintains hepatic glutathione levels such as n-acetyl cysteine. The main isomer and pharmacologically active moiety of silymarin is silibinin (Popp et al., 2020; Saller, Meier & Brignoli, 2001). A meta-analysis of patients poisoned with amatoxins showed that mortality was significantly reduced in patients treated with silibinin compared to those treated without silibinin administration (9.8% with silibinin versus 18.3% with standard treatment; $P < 0.01$) (Saller, Meier & Brignoli, 2001). Silibinin also acts mainly by inhibiting the uptake of amatoxins into hepatocytes via OATP-1B3, like Penicillin G (Popp et al., 2020).

In amatoxin poisonings, silymarin can be administered as 1 g/day oral or as a continuous infusion of 20 mg/kg/day for 6 days after the first 1 hour of 5 mg/kg administration of silibinin intravenously (Mitchell, 2010). It was stated that silibinin (alone or in combination) or N-acetyl-cysteine should be used as the first choice treatment because of the low mortality rates (5.6% and 6.8%, respectively) (Mengs, Pohl & Mitchell, 2012). N-acetyl-cysteine can be given as an alternative if silibinin is not available or in addition to silibinin because of its different mechanism of action (Liu et al., 2020).

Cimetidine inhibits the hepatic microsomal cytochrome P450 enzyme system. For this reason, high dose (200 mg/hour iv) administration in amatoxin poisoning limits the formation of cytotoxic metabolites from amatoxin. Thus, it decreases mitochondrial damage and hepatocellular necrosis (Schneider, Borochoviz & Krenzelok, 1987; Salhanick, Wax & Schneider, 2008).

The combination of methylprednisolone, which inhibits the NTCP transporter, and polymyxin B, which inhibits the OATP 1B3 transporter, has been proposed as a potential antidote due to positive results in animal experiments. However, due to the adverse effects of corticosteroids, more studies are needed before this combination can be applied in humans (Garcia et al., 2019).

Since most of the excretion of amatoxins is via the urinary tract, forced diuresis (for 4 - 5 days with 100 - 200 mL/hr target diuresis) has been recommended to increase their clearance (Jaeger et al., 1993). Hemodialysis and hemoperfusion are used to remove toxins in poisoning. However, their efficacy in amatoxin poisoning is controversial. There was no clinical improvement in patients who underwent hemodialysis and hemoperfusion, and no toxin was found in the dialysis fluids of these patients (Mullins & Horowitz, 2000). The efficacy of hemodialysis, hemoperfusion, and plasmapheresis is limited due to rapid blood clearance (rapid uptake by tissue and low plasma concentrations) of amanitins (Berger & Guss, 2005; Jaeger et al., 1993). However, it has been reported that hemoperfusion may be effective in preventing cerebral edema due to fulminant liver failure (Berger & Guss, 2005). In amatoxin poisoning, hemodialysis is recommended only when acute renal failure develops (Bergis et al., 2012).

In those who develop acute hepatic failure, high-volume plasma exchange has been shown to improve survival without liver transplantation (Larsen et al., 2016). Although plasma exchange transfusions are partially beneficial, their efficacy has been found to be insufficient in controlled studies (Wittebole & Hantson, 2011).

The Prometheus fractionated plasma separation and adsorption (FPSA) system can also prevent liver failure by eliminating amatoxins (Bergis et al., 2012).

Molecular absorbent regeneration system (MARS) is a method that removes albumin-bound toxins with albumin-containing dialysate (Wittebole & Hantson, 2011).

Interruption of the enterohepatic cycle of amatoxins by performing biliary drainage in dogs reduced intestinal amatoxin absorption by more than 70%. Less severe signs of toxicity were observed in dogs with biliary drainage than in dogs without drainage. This could be a promising adjunctive treatment for amatoxin poisoning (Sun et al., 2018).

When treatment is applied in the first 2-3 days of poisoning, mortality decreases to 9% (Saviuc & Flesch, 2003). It was determined that hepatic coma followed the increase in AST and ALT in patients with hepatic failure, and there was a significant correlation between this increase and mortality (Eren et al., 2010).

Liver transplantation should be planned when a complete clinical recovery is not achieved despite all treatments (Enjalbert et al., 2002). It is stated that liver transplantation should be performed in patients accompanied by hepatic encephalopathy with high INR, ammonia and AST levels or patients with metabolic acidosis, persistent hypoglycemia and hypofibrinogenemia (Galler,

Weisenberg & Brasitus, 1992). Ganzert criteria have been developed for liver transplantation in amatoxin poisoning. They stated that at any time between the 3rd and 10th days after eating mushrooms, if the serum creatinine is 106 $\mu\text{mol/L}$ or above with a decrease in the prothrombin index to 25% or less of normal, regardless of the presence of hepatic encephalopathy, urgent liver transplantation is required (Ganzert, Felgenhauer & Zilker, 2005).

Patients should be referred to a more advanced center center to avoid loss of time in suspected amatoxin poisoning. If possible, patients with a diagnosis of amatoxin poisoning should be followed up in healthcare facilities that are experienced in toxicology and have an organ transplant unit (Bernuau, 2004).

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Cognitive Communication Disorders in Alzheimer's Disease

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Introduction

Dementia is a chronic, often progressive syndrome and is the 7th leading cause of death worldwide. In individuals with dementia, cognitive functions are impaired more than would be expected from normal aging processes (Organization, 2017). The result is that dementia significantly reduces people's cognitive functions, activities of daily living, communication skills, and quality of life (Munis, Parlak, & Köse, 2021).

Alzheimer's disease (AD) constitutes 60–70% of the etiology of dementia (Organization, 2017). Although AD first affects the memory, it is a neurodegenerative disease in which cognitive functions such as attention and visual-spatial functions are increasingly impaired (Parlak, Altan, & Saylam, 2022; Parlak, Babademez, Tokgöz, Bizpınar, & Saylam, 2022). Later symptoms can thus include problems with language, mood swings, disorientation (including easily getting lost), self-neglect, loss of motivation, and behavioral issues. It is estimated that AD will become a worldwide public health problem that directly affects 100 million people by 2050. It has been stated that more than five million Americans have AD and that this number may increase to 16 million by 2050 (A. s. Association, 2018; Cao et al., 2020). According to the 2012 Alzheimer Europe Report, 331.512 individuals are currently suffering with AD in Turkey (Özbabalık & Hussein, 2017). Globally, the number of individuals with AD is expected to increase rapidly as the elderly population increases.

Language disorders appear early in AD and worsen as the disease progresses. As the disease progresses and the cognitive functions of individuals deteriorate, the effects on their communication skills increase and their independence level decreases (Parlak, Tokgöz, Bizpınar, Saylam, & Köse, 2022). That people with AD become increasingly dependent on others reduces the quality of life of their caregivers, along with increasing the burdens on families and health systems (KA Bayles & Tomoeda, 2007). In this study, the cognitive communication disorders seen in AD were examined.

Cognition

Cognition is a set of abilities that enable many mental processes and functions to work together and is a general term that refers to our information processing systems and stored knowledge (Baş B, 2022; Kathryn Bayles, McCullough, & Tomoeda, 2018). All situations that enable us to perceive and interpret patterns, to achieve goals, and to change our behaviors to ensure survival are realized through these cognitive functions. Cognitive functions play an important role in many daily living activities, such as chatting with a friend or finding a job, and in adapting an individual to daily life. To conceptualize cognition, we can liken it to a company. The mission of this company is analyzing the senses, detecting regularities in incoming sensory information, remembering, and using experience to guide behavior (Kathryn Bayles et al., 2018; KA Bayles & Tomoeda, 2007).

Cognitive functions are generally evaluated with cognitive screening tests, such as Mini-Mental State Examination and Addenbrooke Cognitive Examination. Cognitive functions are examined in these tests in five sections: orientation, attention, executive functions, memory, visuospatial functions and language. Cognitive functions and their subsections are shown in Figure 1 (Baş B, 2022; Chung, Pollock, Campbell, Durward, & Hagen, 2013; Diamond, 2013; Hoffmann, Bennett, Koh, & McKenna, 2010; Karakaş & İrkeç, 2008; Molloy & Standish, 1997; Noone, 2015; Sohlberg & Mateer, 2001; Squire, 2004).

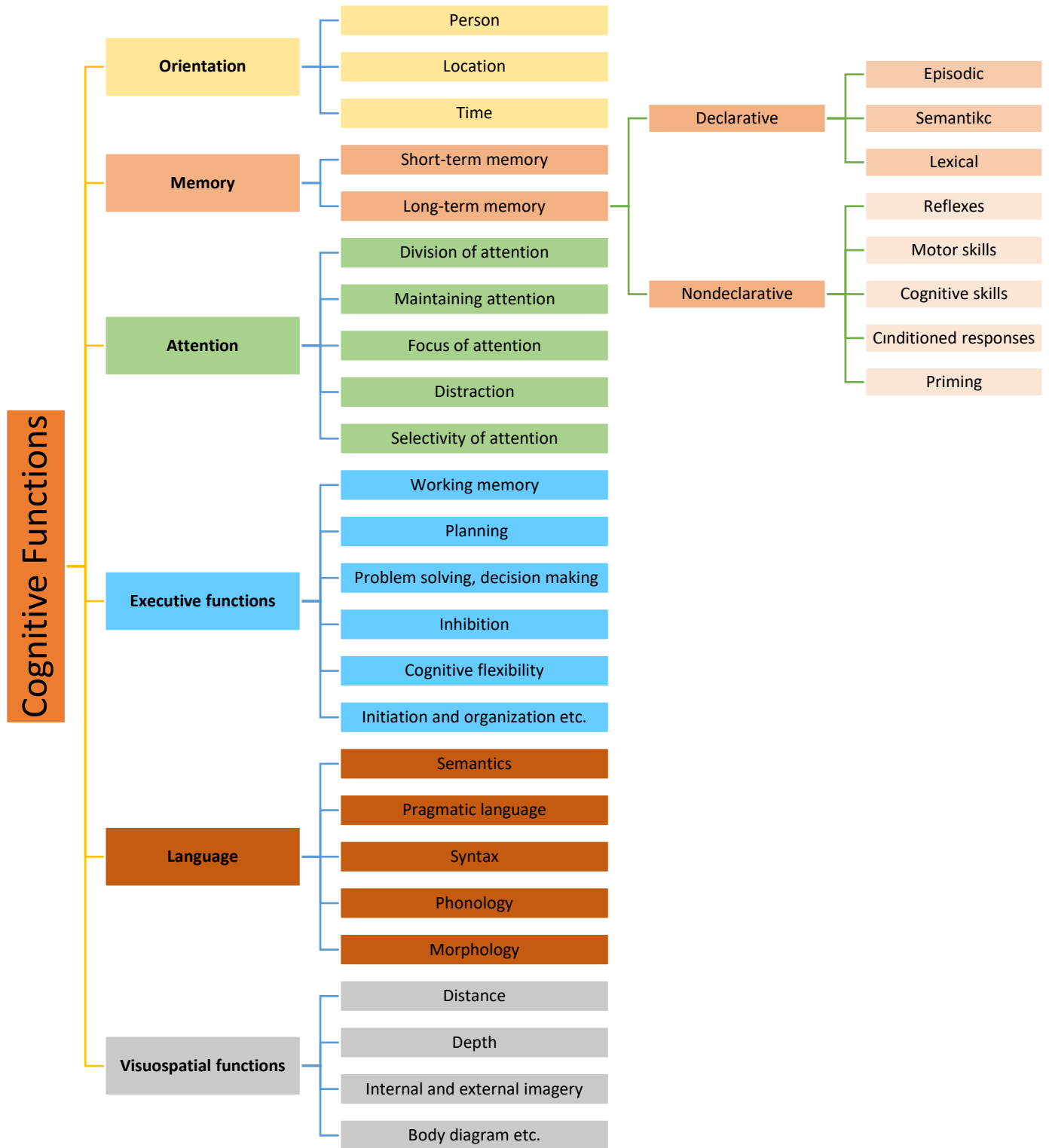


Figure 1: Cognitive functions and their subsections (Baş B, 2022; Chung et al., 2013; Diamond, 2013; T. Hoffmann et al., 2010; Karakaş & İrkeç, 2008; Molloy & Standish, 1997; Noone, 2015; Sohlberg & Mateer, 2001; Squire, 2004)

Communication

Communication is the sharing of information through a system of symbols. Communication starts with the formation of an idea in the speaker's mind. The speaker encodes the message he or she wants to convey in some way; in other words, he or she makes it communicable with the rules of whichever communication tool he or she will use. The receiver of the message analyzes it and makes sense of it (decoding). There are types of communication: verbal communication and nonverbal communication. People convey their thoughts, ideas, and feelings using spoken or written words in verbal communication. Nonverbal communication involves a variety of techniques, including body language, which includes facial emotions, gestures, and more. Communication types are shown in Figure 2 (Hans & Hans, 2015; Liu, 2016; Rocci & de Saussure, 2016).

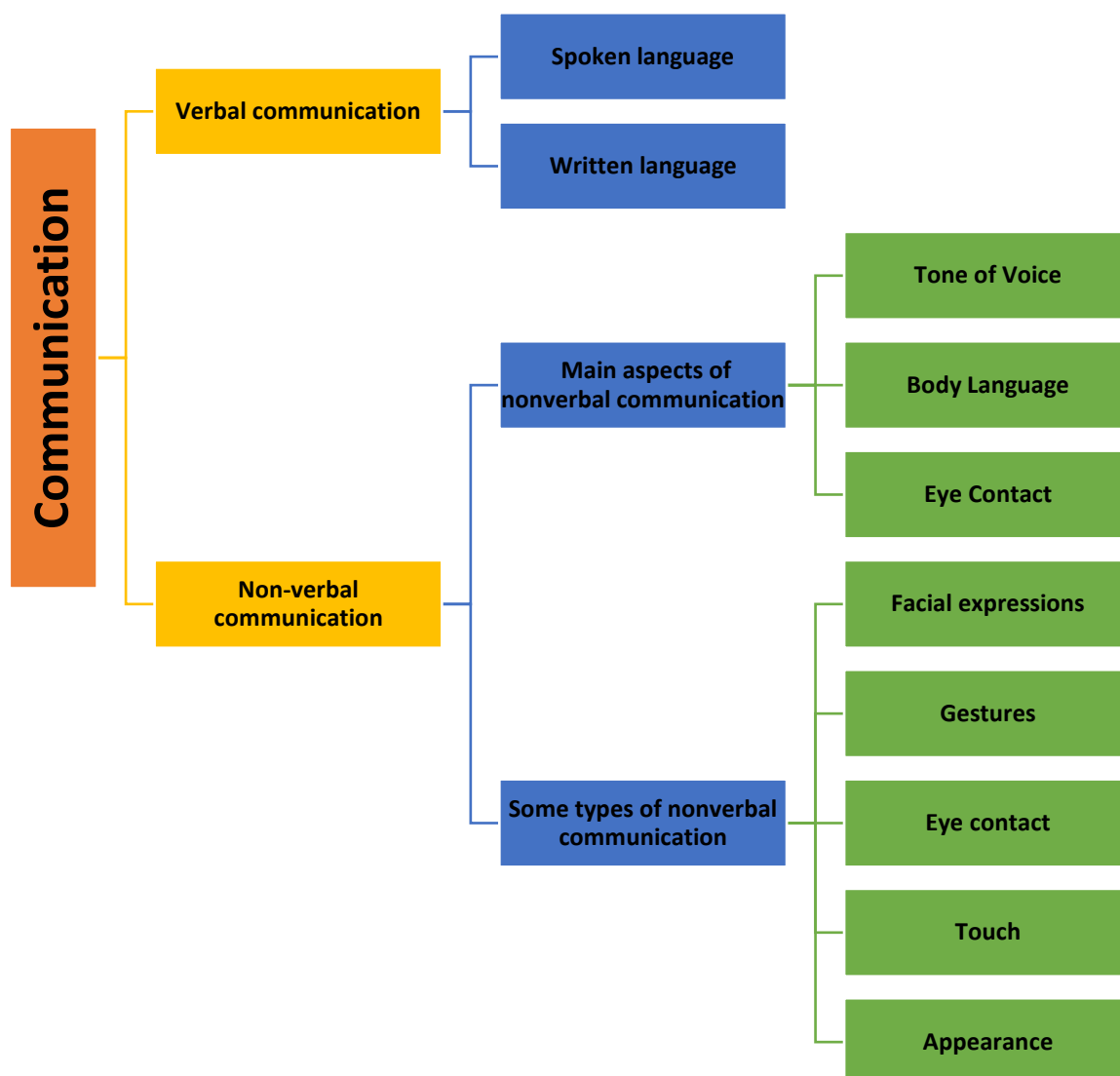


Figure 2: Communication types(Hans & Hans, 2015; Liu, 2016; Rocci & de Saussure, 2016).

To communicate verbally or non-verbally, an individual must have an idea to share and a system of symbols with which to express that idea. Both verbal and nonverbal communication are impaired in AD because both require cognitive processes to share and interpret information. The pathophysiological changes in AD interrupt the information processing process. Because verbal communication requires the conscious sharing of ideas through language, it is typically the most affected in AD (Kathryn Bayles et al., 2018).

Cognitive Communication Disorder

Cognitive communication disorder is difficulty in any aspect of communication skills as a result of the influence on cognitive functions (A. S.-L.-H. Association, 2022c). The most common etiologies of cognitive communication disorders are dementia (especially AD), stroke, brain tumor, traumatic brain injury (TBI), and right brain injury. Cognitive communication disorders have negative effects such as difficulties in communicating people's needs and feelings and reducing their independence. Some potential effects of cognitive communication disorders are shown in Table 1 (A. S.-L.-H. Association, 2022c).

Table 1: *Impacts of cognitive communication disorder (A. S.-L.-H. Association, 2022c).*

Impacts of Cognitive Communication Disorder
Reduced awareness of and ability to initiate and effectively communicate one's needs
Risk of injury due to inability to communicate and/or anticipate the consequences of one's own actions in an emergency
Reduced memory, judgment, and ability to initiate and effectively modify routine information exchange
Reduced social communication skills and/or ability to manage one's emotions, often resulting in loss of relationships
Impairment of ability to fulfill educational or professional roles, resulting in potential loss of employment
Difficulty effectively performing personal lifestyle management activities (i.e., paying bills)
Decreased awareness of the disorder and its extent (i.e., loss of ability to assess one's own communication effectiveness)
Decreased ability to predict possible outcomes with reasonable decision making and problem solving

How does cognitive communication disorder occur in Alzheimer's disease?

The progressive deterioration in cognitive functions in individuals with AD leads to cognitive-communication disorder, preventing communication. The production and understanding of language are linked to information stored in the memory. When we think of the simple act of

naming an object, for example, a carrot. we first need to perceive the characteristics of the carrot. For that recognition to occur, a match must be made with the information in our long-term memory. After that, the intention to say the name of the object is created. The linguistic representations of objects are part of the long-term lexical memory and must be retrieved and brought into consciousness. Maybe you can't be sure of what a carrot looks like, and therefore can't be sure whether the object at hand is a turnip, radish, or carrot. In this case, a decision must be made whether to indicate your uncertainty. A motor plan should be created to express uncertainty about the object's name or properties. Therefore, the simple object naming actions are perception, access to long-term memory, association, recognition, lexical recall, decision-making, motor planning, and self-monitoring (Kathryn Bayles et al., 2018; Klein & Mancinelli, 2021; Szatloczki, Hoffmann, Vincze, Kalman, & Pakaski, 2015).

Since the information processing abilities of the explicit and working memory systems of individuals with AD are affected, they have difficulty in producing verbal information (6, 10, 11). In people with AD, information processing is impaired due to deficiencies in the cognitive processes of perception, recognition, attention, and memory, which causes patients to have difficulties in understanding language (MacDonald, Almor, Henderson, Kempler, & Andersen, 2001).

Cognitive Communication Disorders in Different Stages of Alzheimer Disease

The Clinical Dementia Rating (CDR) scale is frequently used in the clinic for staging AD (Gürvit, 2004; Parlak, Babademez, et al., 2022). The CDR is calculated by testing six different cognitive and behavioral domains: orientation, memory, judgment and problem solving, home and hobbies performance, community affairs, and personal care. The CDR is based on a scale of 0–3: no dementia (CDR = 0), questionable dementia (CDR = 0.5), mild dementia (CDR = 1), moderate dementia (CDR = 2), and severe dementia (CDR = 3) (Figure 2) (Hughes, Berg, Danziger, Coben, & Martin, 1982). In this article, cognitive communication disorders in mild, moderate and advanced stages of AD were examined. Impairments in cognitive functions according to Alzheimer's disease stages are shown in Figure 3 (Kathryn Bayles et al., 2018).

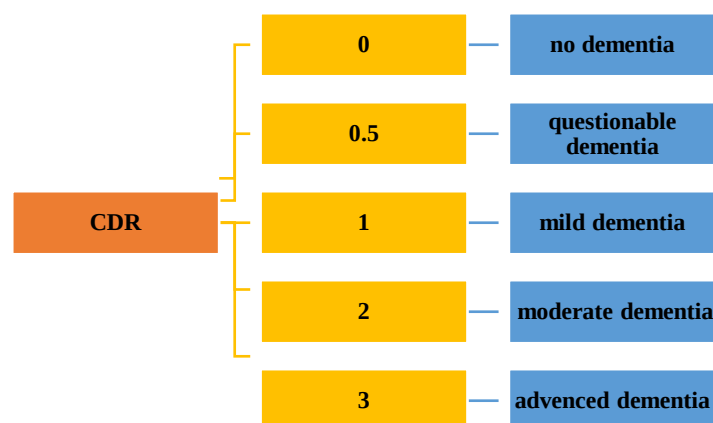


Figure 2: Clinical Dementia Rating Scale (Hughes et al., 1982).

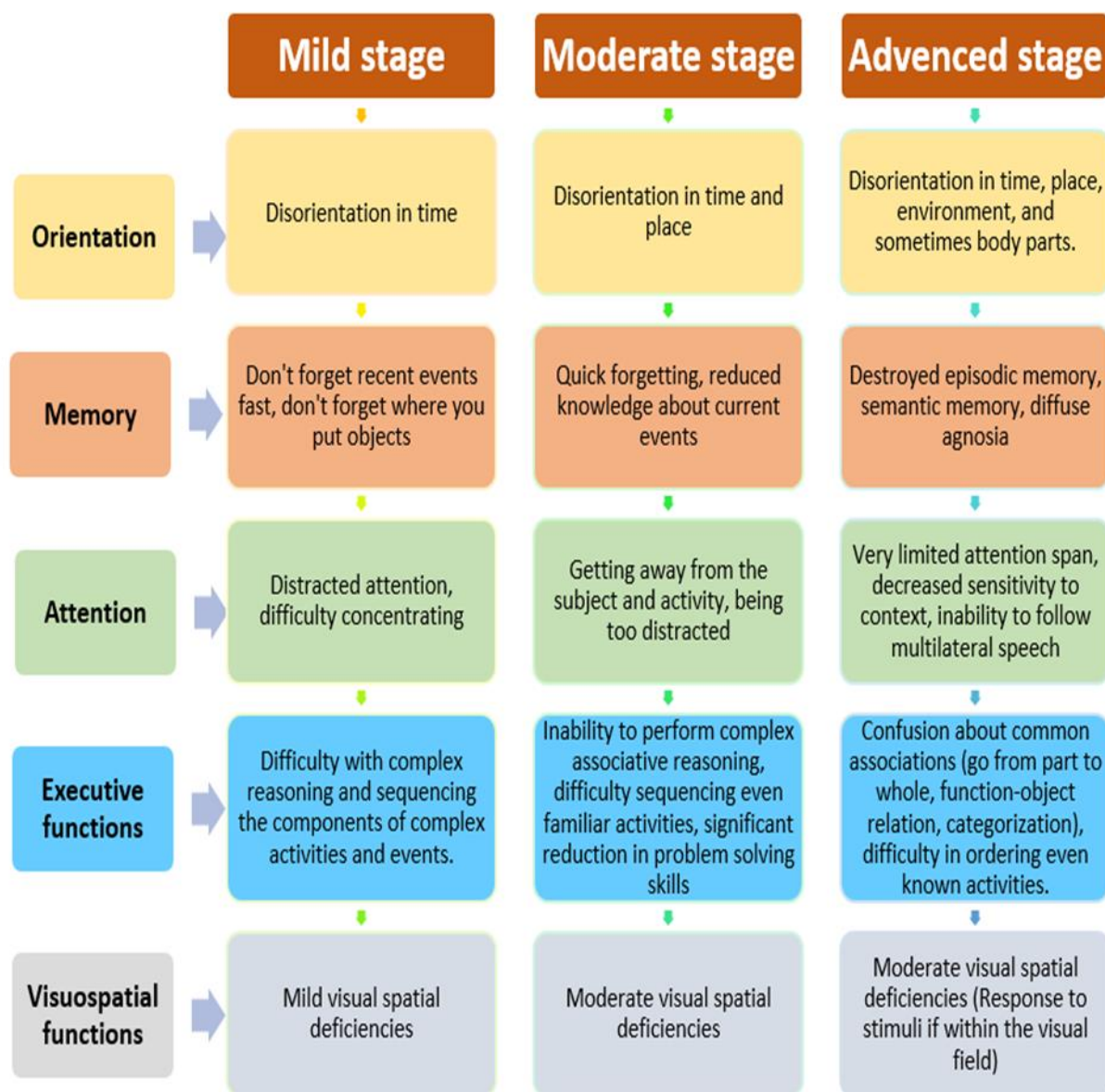


Figure 3: Impairments in cognitive functions according to Alzheimer's disease stages (Kathryn Bayles et al., 2018).

Cognitive Communication Disorders in Mild Alzheimer's Disease

In mild-stage AD, speech is fluent, and the patients can understand most of what they read and hear, but they quickly forget recently acquired information. While the semantic and pragmatic components of the language are affected, phonology and syntax are generally better preserved over the course of the disease. During spontaneous speech, early-stage AD patients usually have longer hesitations and pauses and slower speaking rates (Kathryn Bayles et al., 2018; Gayraud, Lee, & Barkat-Defradas, 2011; I. Hoffmann et al., 2010; Jarrold et al., 2014; Satt, Hoory, König, Aalten, & Robert, 2014). However, their speech is fluent, without dysarthria or articulation errors (Weiner, Neubecker, Bret, & Hynan, 2008).

Grammar is preserved in speech, but there is a marked increase in the number of “empty words” (e.g., “thing” and “it”) in the content of the language (Blanken, Dittmann, Haas, & Wallesch, 1987; Irigaray, 1967). Because early-stage AD patients often forget what they have just

heard or thought, their oral discourse includes more sentence fragments and repetitiveness. Their speech is less consistent than that of their healthy peers (K. A. Bayles, Tomoeda, Kaszniak, Stern, & Eagens, 1985; K. A. Bayles, Tomoeda, & Trosset, 1992). Dysnomia during speech is common, and a naming error is often semantically related to the target word (e.g., “sharp” for “knife”) (K. A. Bayles & Tomoeda, 1983). Their performance on verbal fluency tests shows significant deficiencies compared to that of their healthy peers (Araujo et al., 2011; Henry, Crawford, & Phillips, 2004). In addition, one study found that the most frequently reported symptom in individuals with mild AD was difficulty naming things (K. A. Bayles & Tomoeda, 1991).

Individuals with mild AD can identify words and describe pictures but have difficulty following the context of speech. However, they may have the ability to follow three-step instructions. Reading and writing skills are generally preserved at this stage, but typos are common (Kathryn Bayles et al., 2018; Klein & Mancinelli, 2021; Szatloczki et al., 2015).

The written language of individuals with AD is more affected than their spoken language, and their written discourse contains more errors than that of their healthy peers (Groves-Wright, Neils-Strunjas, Burnett, & O'Neill, 2004). Grammar and spelling errors are common in written language. Although they generally understand what they hear and read, they quickly forget this (K. A. Bayles et al., 1992). They can answer most questions and define words (28), but they often miss the point of a joke, which can be mistaken for sarcasm (Kathryn Bayles et al., 2018).

Cognitive Communication Disorders in Moderate Alzheimer's Disease

People with moderate stage AD are fluent in speech but usually speak more slowly and are more hesitant to speak. Their speech is filled with more frequently repeated words and longer pauses (Gayraud et al., 2011). Intellectual repetitions are common in their speech when they are asked to describe a picture or object. The morphology of the language usually remains intact, but the content is significantly affected (K. A. Bayles et al., 1992).

The oral expressions of individuals with middle-stage AD contain fewer nouns than verbs (Blanken et al., 1987). Vocabulary loss and deterioration in conceptual knowledge are observed to be more common in their performance on vocabulary tests (K. A. Bayles et al., 1992). Their ability to create and name examples of things falling under a certain category is significantly impaired compared to that of their healthy peers. Written languages also contain many errors.

Individuals with middle-stage AD exhibit a decrease in understanding both written and spoken language (K. A. Bayles et al., 1992; Rochon, Waters, & Caplan, 2000). The mechanics of reading are preserved at this stage, although most of them are at the level of words and expressions. However, reading comprehension is impaired, and what they understand from what they read is quickly forgotten. They are largely insensitive to context and may miss the main points of jokes. They also tend to interpret figurative language literally (Kathryn Bayles et al., 2018).

Individuals with moderate AD produce significantly less opinionated discourses than those with mild AD do. During conversations, they often forget their intentions. This leads to many half-sentence fragments. In addition, they lack self-monitoring and self-correction. They are usually able to name items when given naming tasks but have difficulty participating in the conversation. Their verbal outputs tend to be less significant. They usually understand two-step instructions, but they have difficulty understanding three-step instructions. As in the early stage, they can write words for dictation, but they may have difficulty writing long phrases (Kathryn Bayles et al., 2018; Klein & Mancinelli, 2021; Szatloczki et al., 2015).

Cognitive Communication Disorders in Advanced Alzheimer's Disease

In advanced-stage AD, which is the last stage of the disease, due to cognitive destruction, the patients' language outputs are greatly reduced, and their speech is often meaningless. There are significant differences in their ability to communicate with other individuals with advanced AD. Some of them cannot speak at all, while others produce meaningless language. In addition, some may have limited functional vocabulary. Most of the remaining language outputs are either common social expressions or nonsensical words. Many of them are still able to follow simple one-step instructions, but all of them can no longer follow instructions with more than one step. Some can contribute to speech, say their names, and maintain the characteristics of social language (KA Bayles & Tomoeda, 2007; KIM & BAYLES, 2007). However, almost all of them cannot express themselves in writing. Although some can read single words aloud, their reading comprehension is severely impaired (KA Bayles & Tomoeda, 2007). Cognitive communication disorders according to Alzheimer's disease stages are shown in Figure 4 (Kathryn Bayles et al., 2018; Klein & Mancinelli, 2021; Szatloczki et al., 2015).

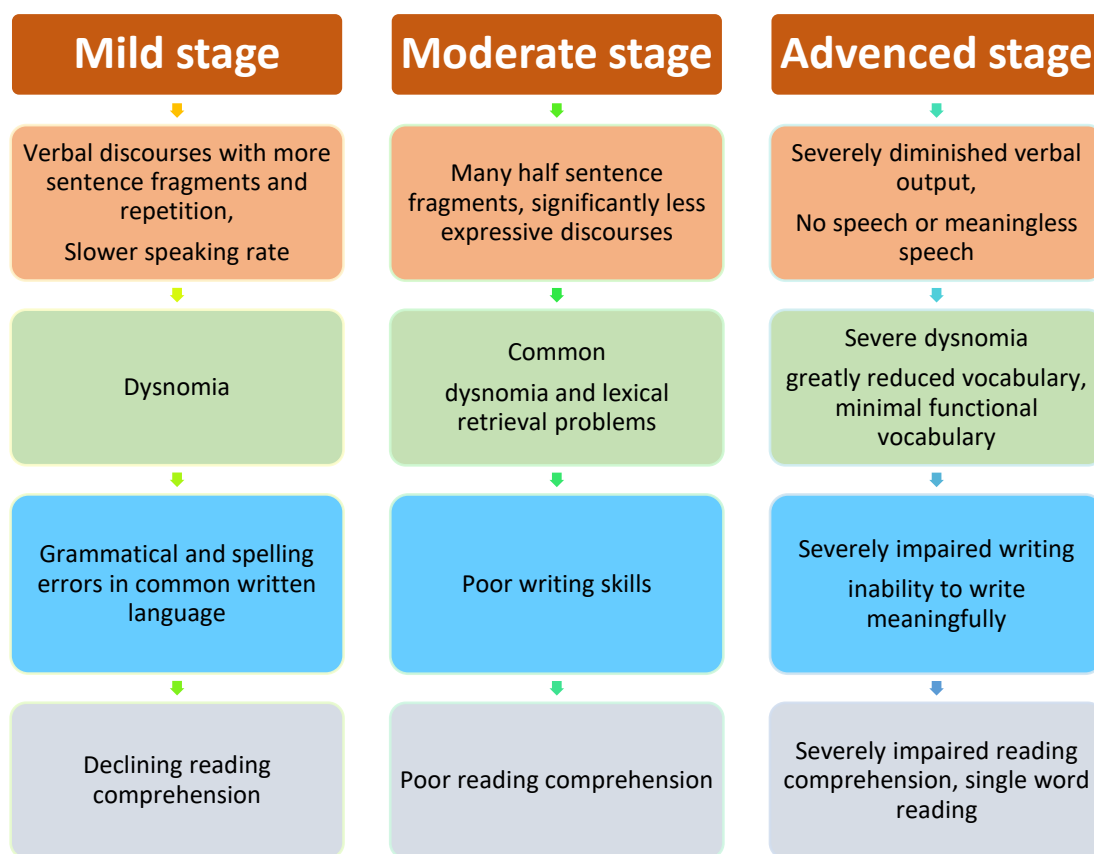


Figure 4: Cognitive communication disorders according to Alzheimer's disease stages (Kathryn Bayles et al., 2018; Klein & Mancinelli, 2021; Szatloczki et al., 2015)

Conclusion

Attention, orientation, executive functions, visuospatial functions, memory, and language are our basic cognitive functions. The most common cause of dementia is Alzheimer's disease, and progressive deterioration in cognitive functions occurs during the disease process. Due to the relationship between communication and cognition, patients' communication skills are affected

even in the disease's early stages. Therefore, cognitive communication disorders are accepted as an important early symptom of Alzheimer's disease, which is a neurodegenerative disease, and communication ability gradually deteriorates over the course of the disease.

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Lumbar Disc Degeneration-Lumbar Disc Hernia

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Introduction

The human lumbar spine, like the entire spine, consists of thin, compressible, fibro-cartilaginous intervertebral discs to support upright posture and allow a wide range of motion. With age, the discs begin to degenerate by losing their water. In certain circumstances, disc damage can accelerate these changes, drastically alter the load-bearing properties of the disc, and lead to significant disc degeneration. In severe cases, this degeneration can cause significant morbidity, including pain and neurological symptoms.

To begin with, disc degeneration as a biochemical phenomenon associated with aging and the accompanying radiological changes differ from disc disease as a symptomatic clinical entity. In fact, radiographic signs of disc degeneration are very common in the general population, even if they are not symptomatic. In the literature, magnetic resonance imaging (MRI) of 67 asymptomatic patients was found in the literature by Boden et al. showed significant abnormalities consistent with its degeneration (Boden et al, 1990). Paaanen et al. in their study, they found abnormal lumbar discs on MRI in 35% of asymptomatic control patients (Paaanen et al, 1989). Disc degeneration and lumbar disc herniation are very common in the community, with or without symptoms.

Symptoms associated with disc disease are often nonspecific; therefore, they do not provide much diagnostic benefit over symptoms. The most common symptom directly related to lumbar disc disease is low back pain, usually without radiculopathy. It should be noted that because back pain is such a common symptom, the presence of back pain is not in itself a diagnosis of disc disease and is not necessarily associated with evidence of disc degeneration on imaging in a particular patient.

Diagnosis

As in all patients, a detailed history should be taken and a comprehensive history should be taken including the onset of symptoms, duration and severity. The symptoms associated with the disease should be identified by identifying aggravating and extenuating causes. It is also important to evaluate the patient's social history in terms of contributing to the patient's treatment. Fever, night sweats, night pain, pain that does not go away with position changes, involuntary weight loss, use of anticoagulants, immunocompromised status, active/previous malignancy, age over 60 years, long-term corticosteroid use, low back pain accompanying severe trauma history are red flag symptoms (Downie's). A et al, 2008, Henschke, Maher & Refshauge, 2008:). Additional investigations should be considered in patients with multiple red flag symptoms. In these patients, symptoms indicating emergencies such as cauda equina syndrome should be sought.

Although not every patient with lumbar disc degeneration shows symptoms, the most common complaint of patients with disc disease is low back pain, usually radiating to the sacroiliac joints (Madigan, 2009). It results in increased sensitivity to inflammatory mediators such as prostaglandin E2 and interleukin-6-8 in the vascularized granulation tissue of annular tears of diseased discs, which affects the severity of pain (Peng, 2006). Pain is usually exacerbated by positions that increase axial loading on the discs and increase intradiscal pressure, such as flexion, standing, and sitting.

Patients who experience protrusion or herniation of the nucleus as a result of degenerative disc disease may develop symptoms of nerve root compression or, in rare cases, neurogenic claudication. The anatomy of the posterior longitudinal ligament is stronger in the midline but weaker laterally. This may facilitate posterolateral herniation and cause compression in the nerve roots (13 Peng 2006). The nerve root affected by the posterolateral herniation will usually be the root that emerges from under the pedicle of the caudal vertebra. These patients may experience symptoms associated with nerve root compression, including dermatomal sciatic pain, distal paresthesias, and myotomal motor weakness.

Physical examination

Since the differential diagnosis of low back pain is wide, it is important to examine not only the waist but also the abdomen and hips. On examination, the patient's gait and standing should be observed. Visual inspection of the lumbar region; and the greater trochanters of the femurs, as well as the spinous processes and paraspinal muscles are important. Spinal range of motion should be evaluated while the patient is standing, especially in the anterior-posterior plane. Patients with pain from disc pathology will typically experience more pain when the axial loading of the discs is greatest and in positions that increase stretching of the lumbar nerve roots. Lower extremity strength, reflexes, and sensation should be tested. Motor and sensory defects of certain muscle groups may indicate the level affected.

Straight leg raise or Lasègue tests are applied for radicular symptoms caused by nerve root compression caused by the lumbar disc. With the patient lying on his back, the affected leg is straightened and slowly passively raised. The test is considered positive if the patient's sciatic pain recurs between 30 and 70 degrees. A meta-analysis by Deville et al. in 2000 revealed the sensitivity and specificity of straight leg lift to be 91% and 26%, respectively, in detecting lumbar disc herniation (Deville, 2000). The onset of symptoms with contralateral leg elevation - the so-called cross straight leg raise or Fajersztajn's cross sciatic sign, while not particularly sensitive (29%) is much more specific (88%) (Deville, 2000).

Treatment

Non-Surgical Treatments

The initial treatment for acute low back pain, including those caused by lumbar disc disease, is usually nonsurgical. The primary treatment for acute and subacute low back pain in adults is patient education, activity modification to limit disc loading, nonsteroidal anti-inflammatory drugs (NSAIDs), and gradual return to physical activity. However, some patients may find relief with heat therapy. Muscle relaxants can provide relief, but their side effects should be kept in mind. Limited dose and limited duration opioid use may be considered in a selected group of patients with severe acute pain that cannot be adequately controlled with NSAIDs and acetaminophen (Goertz, 2012, Modic, 1984).

The results of epidural steroid injections for acute low back pain in the literature are not consistent with each other. There is some evidence that they may provide short-term subjective improvement in low back pain, but this is not consistent with a functional improvement or change

in surgical planning. Epidural injections are not recommended in patients without symptoms of radiculopathy. There is no indication for oral steroid use in the treatment of uncomplicated acute low back pain (Engers, 2008, Heymans, 2008).

Physical therapy applications such as acupuncture, spinal manipulation, transcutaneous electrical nerve stimulation or lumbar traction may also play an active role in pain. However, surgical intervention may be considered for patients with prolonged symptoms or progressive or particularly motor weakness of pain and neurological symptoms (Engers, 2008).

Surgical Treatments

Since most cases of low back pain (>85% in 6 weeks) (including those with disc herniation) are self-limiting and will resolve with non-surgical treatment in the first place, they can be followed up with non-surgical treatments, except in emergencies. It is important to select patients who will require surgical intervention. (1) patients must be symptomatic; (2) symptoms must match examination and imaging findings; (3) symptoms must be resistant to a long course of appropriate non-surgical treatment (5-8 weeks); (4) there should be a reasonable expectation that surgery will improve the patient's symptoms and functional status, (5) patients with persistent, debilitating low back pain due to painful disc herniations or disc degeneration and who do not respond to nonsurgical treatment may be considered for surgery. In patients with one or two levels of degenerative disc disease and chronic low back pain for at least 2 years, research has shown better benefit of lumbar spinal fusion versus intensive physical therapy (Engers, 2005).

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Effects of Melatonin on Physiological Functions

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INTRODUCTION

Melatonin is a derivative of the serotonin molecule produced in our body as a result of the metabolism of the amino acid tryptophan. The serotonin molecule, which is widely produced in the brain, is converted into melatonin in the pineal gland and given to the blood. Studies have shown that the hormone melatonin plays a very important role in protecting and improving body health. Melatonin is responsible for circadian regulation and sleeps control, which is encountered in the organism at the highest concentration during the night (Tamura et al., 2009). With ageing, there is a significant decrease in melatonin production among the elderly, which may decrease sleep quality (Kennaway et al., 2002). In humans, as in other mammals, pineal melatonin is the nocturnal hormone. It shows a circadian rhythm in the blood, and nighttime values are considerably higher than daytime levels (Waldhauser et al., 1984; Ilnertova, 1988). Melatonin has many important functions in metabolism. Melatonin is also known to affect the depth and quality of sleep. Accordingly, it has been determined that people who secrete enough melatonin have a more restful and relaxing sleep (Tsuno et al. 2005). In this way, melatonin; is also a regulatory factor for cardiovascular health, blood pressure, immune system functions and hormone levels (Pieri 2013). Normally, various oxidizing and harmful substances are also produced as a result of chemical reactions occurring in the physiological order of our body. These substances are called reactive oxygen radicals. Radicals play a critical role in the development of various diseases such as ageing, cardiovascular diseases, infections and cancer (Reiter et al. 2000). Melatonin; protects the health of the body by cleaning and removing the radicals produced in the body during the night

(Maestroni et al.1999). Studies show that melatonin secretion alleviates the effects of depression, is beneficial in relieving reflux and supports eye health (Konturek et al. 2007). Melatonin has also been reported to trigger growth hormone production from the pituitary gland during secretion. Growth hormone plays a critical role in the regulation of body metabolism. There are studies that the secretion of melatonin prevents the development of diabetes. It has been observed that the level of fat and cholesterol in the blood decreases in proportion to its production, therefore it plays a supporting role in heart health (Arendt 2000).

DISCUSSION

1. Effect of melatonin on the reproductive system

Many studies have shown that melatonin secretion affects the reproductive system (Öztekin et al., 2006; Öztürk et al., 2003). Although exogenous melatonin varies according to species, age, dose and application time, it modifies reproduction (Turgut et al., 2002).

In seasonally breeding hamsters, long-term darkness inhibits reproduction due to more melatonin secretion, causing testicular regression in males and anoestrus in females. Although humans do not show seasonal reproduction, epidemiological studies reveal that the pregnancy and

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birth rates show the seasonal distribution in different geographical areas. In regions with long winters, activation of the hypothalamic-gonadal system and pregnancy are lower than in summer (Ölmez et al., 2000). The pineal gland is an upper centre that regulates the endocrinological activity in the organism through the pituitary, thyroid, adrenal gland and gonads. Therefore, it is closely related to events related to dysfunctions in endocrine organs (Turgut et al., 2002).

When the melatonin level drops, the hypothalamogonadal system is activated. During childhood and adolescence, melatonin synthesis gradually declines. Serum melatonin concentration was found to be high in hypothalamic amonites. With rapid and continuous (if prolonged) exercise, the serum melatonin concentration rises, which can cause ammonia. Melatonin inhibits gonadotropin-releasing hormone (GnRH) in experimental animals. Similar data in humans have not yet been fully elucidated (Brzezinski, 1997). It has been shown that when 300 mg of oral melatonin is given to young women for 4 months, it inhibits LH release and ovulation, and progestin administration increases this effect (Olcese, 2020). It can also directly affect the functions of the ovaries. The presence of melatonin receptors in the granulosa cell membrane has been demonstrated (Voordouw et al., 1992).

In a study of mice, mice administered melatonin was able to maintain sexual interest and performance similar to much younger mice; It has been reported that both male and female mice show regeneration of the sex organs (Gunn et al., 2016)

2. Effect of melatonin on the immune system

The first finding of the immunological role of melatonin belongs to Mills et al. (2005). Carrillo et al. reported that suppression of immune functions as a result of soft tissue trauma and hemorrhagic shock in mice was reversed with melatonin and that chronic melatonin treatment increased leukocyte natural killer (NK) activity in humans (Carrillo-Vico et al. 2013).

Melatonin increases the immune response. When applied in immunodeficiency states, it leads to significant immune activation. Acute stress, which is known to have negative effects on immune parameters, and immunodeficiency conditions caused by the application of immunosuppressive pharmacological agents can be controlled with melatonin (Ölmez et al., 2000).

Innate immunity can be modulated by the exogenous administration of melatonin. NK cell activity is reduced following pinealectomy in mice. The presence of melatonin in the bone marrow indicates its modulating effect on the growth of NK cells and monocytes. An increase in both NK cells and monocyte counts was observed 7-14 days after the administration of exogenous melatonin to young male mice. In addition, melatonin mediates macrophage production and function. The production of cytokines such as TNF- α and IL-1 β , known as macrophages, can be altered by the external administration of melatonin (Hotchkiss et al., 2002).

The immunomodulatory effect of melatonin is mediated by opioid peptides, lymphokines and pituitary hormones originating from T-helper lymphocytes. It also causes a significant increase in interleukin-2 (especially the production of T-lymphocytes, which play an important role in the body's fight against cancer) and γ -interferon levels.

Melatonin also increases the *in vivo* antibody response (IgM and IgG) (Lissoni et al., 1994). Since the inhibitory effect of melatonin on the immune response is related to the antioxidant property of the molecule, it has been suggested that it may be useful in organ transplantation. The lack of toxicity also supports that this agent can be used safely in transplantation (Reiter et al., 1999).

Another effect of melatonin on the immune system is its anti-inflammatory effect. Some studies suggest that melatonin may affect diseases that may contain viruses and be contagious, such as HIV, bacterial infections, and cancer (Maestroni, 1999).

3. Effect of melatonin on ageing

The pineal gland synthesizes melatonin, a powerful free radical scavenger and antioxidant agent, and its functions decrease with ageing; This gland has increased the interest of researchers dealing with ageing (Reiter et al., 1998).

Decreased antioxidant capacity and damage caused by free radicals are also mentioned among the causes of anatomical and functional degeneration occurring in organs during the ageing process, and it is reported that the reason is melatonin, which decreases with age (Mollaoğlu et al., 2005; Kılıç et al. 2020). Because melatonin is thought to have a stimulating effect on antioxidant enzymes, reducing lipid peroxidation and protecting brain tissue from oxidative damage. Increased free radicals due to decreased melatonin have been blamed for many of the neurodegenerative damages that occur in the brain (Kerman et al., 2005).

Many scientific articles report that plasma melatonin concentration in humans decreases with age (Touitou, 2001; Turgut et al., 2002). With ageing, there is a decrease in the synthesis and release of melatonin. Parallel to this, the circadian rhythm of melatonin is disrupted. According to Touitou (2001), the plasma melatonin concentration measured in elderly people is approximately 40-50% lower than the normal value. In experimental animals, it was determined that the circadian rhythm deteriorates as the animal ages, and the serum melatonin levels of day and night become almost equal (Turgut et al., 2002).

4. Effect of melatonin on sleep

Sleep is indispensable for humans. The key mechanism in sleep regulation is the light-dark cycle, and exposure to light leads to stimulation of a neural pathway that extends from the retina to the hypothalamic area in the brain. The suprachiasmatic nucleus, located in the hypothalamic area, works like the "biological clock" responsible for regulating activities that affect the whole body, by initiating signals that control hormones, body temperature, and the feeling of sleep or wakefulness to other areas of the brain (Macchi and Bruce, 2004). Until the dark hours come, the release of sleep-related melatonin-like hormones is suppressed due to the inactivity of the pineal gland. When the sun's rays disappear, the pineal gland is stimulated and melatonin production begins, as the suppressive signals that prevent the release of melatonin created by the suprachiasmatic nucleus disappear. As melatonin levels increase, fewer stimuli are perceived and the feeling of sleep increases (Arendt, 2000; Arendt et al., 2005).

In a study by Garfunkel et al.; It has been reported that melatonin deficiency has an important role in the frequency of sleep disorders in elderly people, that there is no change in total sleep time with controlled-release melatonin at a dose of 2 mg/night, and sleep quality effectively improves in these patients with sleep disorders (Garfunkel et al., 1990). In another study, it was reported that melatonin supplementation given to patients with melatonin deficiency and sleep disorder may be beneficial in falling asleep (starting sleep) and in maintaining sleep (Haimov et al., 1995).

Riemann et al. (2002); showed that the nocturnal melatonin concentration decreased in patients with insomnia. Mac Farage et al., (1991); investigated the effect of using high-dose melatonin (75 mg) in the treatment of chronic insomnia and found that total sleep times increased in the group given melatonin. Low doses of melatonin have also been shown to be effective in treating insomnia. Volunteer patients were given 0.3 mg and 1 mg doses of melatonin, and it was shown that melatonin at both dose levels reduced the time needed to fall asleep (Birdsall, 1996).

Jet lag is a disorder that manifests itself with insomnia and flu-like symptoms that develop due to the deterioration of body rhythm after overseas flight travel. Concomitant concentration and disorientation, a decrease in blood pressure, and blood sugar levels, and changes in energy, alertness and hormone levels can be observed (Rohr and Herold, 2002; Ölmez et al., 2000). It has

been reported that the administration of melatonin minimizes the effects of jet lag in overseas travellers (Birdsall, 1996).

5. Effect of melatonin on cancer

It has been found that melatonin stops cell proliferation in cancerous tissues, inhibits mitotic activity, and has an antiestrogen effect in breast tissue (Macchi and Bruce, 2004). These effects are likened to chemotherapeutic agents. Therefore, melatonin reduces the proliferation of cancer cells, tumour growth and the number of metastases. In addition, melatonin levels were found to be low in patients with prostate and breast cancer (Macchi and Bruce, 2004; Özgüner et al., 1995).

In some studies, it has been reported that pinealectomy increases tumour formation, while melatonin decreases it (Özgüner et al., 1995). Similarly, melatonin has been found to reduce the frequency of chemical-induced tumours in animals. Pinealectomy; It has been determined that while it increases experimental cancer growth and metastasis of melanoma, leukaemia, lung, liver, ovary, pituitary gland, and prostate, high melatonin amount suppresses cancer growth in these tissues (Regelson and Pierpaoli, 1987; Karasket and Fraschini, 1991).

In a study conducted with the addition of melatonin to the treatment in patients receiving immunotherapy and chemotherapy; In addition to the regression in the tumour, a decrease in the severity of side effects and a prolongation of the life span has been reported (Lissoni et al., 1996).

It is reported that the probability of cancer is low in visually impaired people (Coleman and Reiter, 1992; Bartsch et al., 1993). In this statement; It has been shown as a possible mechanism that melatonin is secreted more in visually impaired individuals. In this context, in the studies of Feychting et al. in Sweden and Verkasalo et al. in Finland, it was determined that completely blind women have a lower risk of breast cancer compared to the general female population (Feychting et al., 1998; Verkasalo et al., 1999).

6. Effect of melatonin on the cardiovascular system

Recent findings have shown that the effects of melatonin on the cardiovascular system are receptor and non-receptor mediated. Melatonin causes vasoconstriction in cerebral arteries and vasodilation in peripheral vascular beds. Melatonin levels were found to be low in coronary heart patients with myocardial infarction risk and sudden death risk (Sewerynek, 2002; Dubocovich and Markowska, 2005). Similarly, melatonin levels are low in patients with high LDL cholesterol levels and also in hypertensive patients, and it has been shown that melatonin administration reduces blood pressure (Paulis and Simko, 2007).

7. Effects of melatonin on feeding behaviour

There are different types of studies with conflicting results regarding the effect of melatonin on food intake. While some researchers have shown that melatonin reduces food intake in rats, chickens, hamsters and fish, others have suggested that melatonin has no effect on food intake in rats. However, studies in rodents have reported that food intake increases in response to the exogenous administration of melatonin or its agonists (Angers et al., 2003). It was observed that the blood glucose level, which increased after intravenous glucose administration, decreased again during the sleep period. Melatonin is thought to play an important role in this decrease in blood glucose levels (Rohr and Herold, 2002).

CONCLUSION

The basic duties of the hormone melatonin in the body can be summarized as follows: Melatonin is primarily responsible for the regulation of the body's sleep pattern. Accordingly, thanks to melatonin, whose secretion rate changes in the day-night cycle, the need for sleep occurs at night when daylight is lost. Again, thanks to melatonin, awakening from sleep is provided at certain times of the day through regulated sleep time. It is also known that melatonin secretion suppresses the programmed cell death (apoptosis) mechanism in cells (Yu Q et al. 2000). In this way, it is ensured that the life span of sensitive cells, which are of great importance for the body, especially neurons, which are the basic cellular elements of the nervous system, is prolonged and their loss is prevented. melatonin; helps to slow down the ageing process of the skin by increasing the sleep pattern and quality. Ageing is associated with oxygen radicals and wear of body tissues Melatonin creates an opportunity for tissues to repair themselves during sleep.

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Importance of Milk in Nutrition

Nurhayat Atasoy¹

Introduction

The history of milk begins in the Neolithic Age when people transitioned from nomadic hunting to a settled society. With the establishment of a fixed dwelling, it was possible to collect and domesticate animals for work and food, thereby improving agricultural practices. The first animals to be domesticated were goats and sheep from the Middle East, which made it easier for humans to manage due to their size, robustness, adaptability, behaviour and social nature. For a long time, goats and sheep were a source of food (meat and milk) and clothing (wool). For centuries, milk has become a desirable and valuable source of food; thus, herds were formed and dairy breeds were selected (Yildiz,2010; Barłowska et al., 2011). It is seen that the themes of milk and milk curd are worked in the Babylonian reliefs of the 26th century BC. Again, in the 8th century BC, there are expressions about milk, curd and cheese in Homer's writings (Jain, 1998).

Humans have developed close contact with nature, and milk has become the 1st food source after birth. Breast milk has always been the sole source of food for children. Milk was introduced only after the domestication of animals, starting from goats and sheep about 13000 years ago (Haenlein, 2007; Yildiz, 2010), after nursing, followed by the domestication of cows about 4000 years later (Yıldız, 2010). Therefore, milk has become an indispensable food in human life.

Milk contains more vital nutrients than other foods. The nutritional value of a food is measured by the content of nutrients that the body needs to perform its normal functions. Almost all of the nutrients required by the body in milk have been collected in a sufficient and balanced way. Therefore, it is a foodstuff with superior properties (Gerhardt and Thomas, 2006).

Importance and Properties of Milk

Milk is a secretion of the mammary gland, the physical properties and composition of which vary between species. If a section of the breast is taken and examined; It is seen that a large number of channels, the diameter and number of which vary according to species and race, come to each udder chamber, which is separate and independent in terms of function (Demirci, 2001). Milk is a mixed oil/water emulsion containing fat, protein, lactose, minerals, enzymes, cells, hormones, immunoglobulins and vitamins. (Selvaggi et al., 2014b).

The blood vessels in the breast must supply sufficient blood to the cells and tissues that make up the milk. Because an average of 350-500 litres of blood is required for the formation of 1 litre of milk. It is synthesized in milk alveolar cells. While some components of milk pass directly through the blood, most of them are re-synthesized with the cornerstones in the blood (Metin, 2005). Worldwide, cow's milk is the most consumed, dominating world milk production with 782 million tons in 2013. Therefore, 85% of world milk production comes from cattle, followed by milk from other species such as Buffalo (11%), Goat (2.3%), Sheep (1.4%) and Camel (0.2%)

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(FAO, 2015). However, farms producing sheep milk constitute an important part of agricultural economies in many countries, especially in neighbouring countries to the Mediterranean and the Middle East. The world's largest producer of sheep's milk is China (12.2%) and the leading producers in Europe are Greece (8.7%), followed by Romania (7.2%) and Italy (6.1%) (Barłowska et al., 2011). Sheep milk is slightly less important in Near East and North Africa with 7.5% production and in Sub-Saharan Africa (5.6%) and East and Southeast Asia (3.9%). Milk production from small ruminants such as sheep and goats has increased over the years and new consumer markets are now sought (Selvaggi et al., 2014a). Milk is a good source of minerals such as calcium, phosphorus, magnesium, potassium, and zinc. However, milk with low iron content and iron bioavailability cannot make a significant contribution to the iron requirement in childhood. The mineral content of milk is affected by many conditions such as the physiological condition of the animal, lactation status, environmental and genetic factors, and some processes applied to milk (Baysal, 2004).

Under normal conditions, milk has a light consistency and a homogeneous fluidity. However, in some cases, this appearance may change; creeping, smearing, and the sticky structure may occur. If it shows a very thick consistency, it indicates that the milk has been mixed with colostrum or maybe late lactation milk. Milk has a slightly sweet, pleasant flavour provided by lactose, fat and minerals. The taste and smell of milk with high dry matter are perceived more strongly. The taste and odour in milk are revealed by the effect of some aroma substances. It is known that there are traces of flavour substances such as acetone, acetaldehyde, butyric acid and other free acids in fresh milk. As a result of increased chlorine ions and decreased lactose in breast diseases, milk also has a slightly salty taste. Excess globulin and mineral substance in colostrum give the milk a bitter and salty taste, as seen at the end of lactation (Metin, 2001).

Milk normally has a white or creamy colour. The natural colour of milk is affected by the breed and diet of the dairy animal. Milk is perceived as porcelain white with the effect of colloidal substances such as opaque calcium caseinate and milk fat reflecting light. After the casein is separated, the remaining whey appears in greenish-yellow colour, and in skimmed milk, it appears white with a slight blueish colour (Kırdar, 2001).

Almost all of the essential vitamins for humans are found in milk. Vitamins A, D, E and K are associated with milk fat. It is the carotenoids that give milk fat its yellowish colour and the riboflavin that gives it its fluorescent colour. As milk fat decreases, the content of fat-soluble vitamins also decreases. Unenriched milk has very little vitamin D and K. Milk also contains water-soluble vitamins. It is considered a good source of folate because it contains folate-binding proteins and whey protein that increase absorption (Miller et al., 2000).

Milk fat affects the appearance, taste, flavour and durability of milk. It is also a source of essential fatty acids, fat-soluble vitamins and energy. The oil exists as microscopic globules in the water emulsion. Milk, triglycerides (97-98%), phospholipids (0.2-1.0%), free sterols (0.22-0.41%), free fatty acids, fat-soluble vitamins (A, D, E, K), contains more than 400 different fatty acids and fatty acid derivatives (Miller et al., 2000).

The specific gravity of Cow's Milk is 1.028-1.037 g/cm³, slightly higher than that of water. The reason for this difference; is lactose, protein and minerals found in milk with specific weights varying between 1.6-3.0 g/cm³ (Kırdar, 2001).

Due to the lactose and minerals found in real solution in its composition, milk freezes at a lower temperature, at about -0.55°C, compared to distilled water (Oysun, 1991).

Lactose and soluble minerals, which are in the structure of milk and form a real solution, increase the boiling point. Due to these substances, the boiling point is 100.16 °C (Besler and Ünal, 2006).

DISCUSSION

Minor Components of Milk

Milk is often described as a colloidal suspension containing emulsified fat globules, a heterogeneous family of major and minor proteins, carbohydrates lactose, minerals, vitamins, enzymes (Huppertz and Kelly, 2009) and many minor components with important physiological and metabolic properties. It includes technological roles such as cytokines, nucleotides, peptides, polyamines, enzymes and other bioactive peptides (Haug et al., 2007).

The high content of saturated fatty acids in milk has been reported to be responsible for adverse effects that contribute to heart disease, weight gain and obesity (Insel et al., 2004). However, this is controversial as many milk components support health benefits, including oleic acid, conjugated linoleic acid, omega-3 fatty acids, proteins, vitamins, minerals and bioactive compounds, and various milk proteins and their peptides. -It has been suggested that it has cancer activity (Duarte et al., 2011; Rodrigues et al., 2009).

Salts and Minerals

Milk salts are mainly phosphates, citrates, chlorides, sulfates, carbonates and bicarbonates of sodium, potassium, calcium and magnesium. Since milk contains organic and inorganic salts, the salt level is by no means equivalent to the ash content (Huppertz and Kelly, 2009).

Some elements enter the milk from food, but milking equipment and equipment are important sources of elements such as copper, iron, nickel and zinc. The mineral and vitamin contents of goat and sheep milk are mostly higher than cow's milk (Park et al., 2007). Vitaminler

While milk's fat-soluble vitamins A, D, E, and K are related to the fat ratio of milk, water-soluble vitamin B complex and vitamin C are related to the water phase. Vitamins are variable and therefore their processing can reduce the effective vitamin content in milk. During processing, fat-soluble vitamins are retained by the cream, while water-soluble vitamins remain in skim milk or whey (Michaelidou and Steijns, 2006).

Immune Components

Milk plays an important role in the defence of the mammalian host (Stelwagen et al., 2009). Immunoglobulins (Ig) found in the colostrum and milk of all lactating species provide immunological protection for the offspring against microbial pathogens and toxins. Depending on the species, it can find different immunoglobulins and concentrations. In colostrum, the concentration of immunoglobulins is particularly high, in contrast to IgG, the important class of immunoglobulins in ruminant milk, and the main immunoglobulin IgA found in human milk. Immunoglobulins are transported to mammary secretions via special receptors. In addition to immune globules, both colostrum and milk contain living cells, including neutrophils and macrophages, which secrete some of the immune-related components. These include cytokines and antimicrobial proteins and peptides such as lactoferrin, defensins and cathelicidins. Mammalian epithelial cells themselves contribute to hosting defence by secreting several innate immune effector molecules. A detailed understanding of these proteins and peptides offers great potential to add value to the dairy industry. This is demonstrated by the widespread commercial applications of lactoferrin isolated from bovine milk (Rodrigues et al., 2009).

Bioactive Peptides

Bioactive peptides can be obtained from precursor proteins by enzymatic hydrolysis with digestive enzymes derived from microorganisms or plants, or by fermentation of milk with proteolytic starter cultures (Korhonen and Pihlanto-Leppala, 2006).

Among the bioactive peptides from milk, those with blood-pressure-lowering effects are of particular interest (FitzGerald et al., 2004). Some antihypertensive products based on milk peptides with clinically proven health benefits are commercially available (Lopez-Fandino et al., 2006). Sheep and goat milk proteins are important sources of bioactive ACE-inhibiting peptides and antihypertensive peptides (Park et al., 2007). Goat milk is seen as an attractive area of research as it is less well studied than bovine milk, and also because new peptide angiotensin-converting enzyme (ACE) inhibitors are found in goat milk hydrolysates (Geerlings et al., 2006).

Polyamines

Polyamines consist of flexible polycations that are fully charged under physiological pH conditions. It fulfils several roles in cell metabolism and is necessary for cell growth and proliferation (Löser, 2000; Eliassen et al., 2002; Gugliucci, 2004; Larqué et al., 2007). In addition to participating in DNA, RNA and protein synthesis, the most important function of polyamines is to mediate the interaction of all known hormones and growth factors.

Polyamine requirements that cannot be met by biosynthesis must be met with exogenous polyamines consumed from food (Jeevanandam et al., 1997). It has been suggested that maturation of the gut is maintained by dietary polyamines; therefore, supplementation may be beneficial in formula-fed infants.

Moreover, polyamines are important for the reliability of enhanced DNA transcription and RNA translation that occur in response to infection and during tissue repair, during intestinal growth after surgery, and in intestinal barrier functions (Gimble and Gimble, 1998).

Hormones

Milk contains varying levels of hormones that have limited nutritional or diagnostic value. However, many studies have been conducted on the physiological roles of hormones in human and bovine milk (Koldovsky and Thornburg, 1987; Grosvenor et al., 1993).

Estrogen concentrations, namely 17 β -estradiol, estrone and estriol concentrations in milk and a few dairy products, have been reported (Wolford and Argoudelis, 1979). Cow's milk fat contained 65% 17 β -estradiol and 80% estrone. The occurrence of estrogen in both butter and skim milk indicates the distribution of these steroids between the milk fat phase and the serum phase. Estrone is found in milk as the predominant estrogen and estrogen concentrations are well known to correlate with the pregnancy and reproductive cycle. Progesterone levels in milk, it is associated with pregnancy and childbirth (Comin et al., 2005).

Cow Milk

When you think of milk, the first thing that comes to mind is cow's milk. Due to the high yield and the length of the lactation period, only cows are used as dairy animals in many countries today (Kırdar, 2010).

Most cows are born just before spring and lactation stop for 8-10 weeks during the winter. This production system is characterized by irregularities in the milk supply to the processor, both in quantity and composition, accompanied by seasonal variations in milk production potential (Auldist et al., 1998). In particular, milk coming from the late hours of the production season (autumn and winter) may have different production characteristics than early and mid-season milk, and some products cannot be produced at all (Lucey, 1996).

Sheep Milk

The nutritional value of sheep milk is higher than that of goat and cow milk, with proteins, lipids, minerals and vitamins important for human health and a caloric value corresponding to 5932 kJ/kg (Haenlein, 2001; Kaminarides et al., 2007; Park et al., 2007; Barłowska et al., 2011).

Since approximately 80% of the proteinaceous substances in its composition are made up of casein, it is included in the group of casein milk. Due to its high dry matter, its caloric value is also high. Its colour is whiter than cow's milk. The average lactation period of sheep is 7 months and their milk yield is between 400-700 litres in a lactation period, depending on the breeds (Akçapınar, 2000).

Sheep milk is a rich source of minerals. Calcium, phosphorus, magnesium, zinc, manganese and copper levels are higher in sheep compared to cow's milk, while the reverse is seen for potassium and sodium (Park et al., 2007; Wijesinha-Bettoni and Burlingame, 2013). Calcium and phosphate, which are essential elements for growth and bone maintenance and necessary for newborns, are abundant in sheep's milk (Al-Wabel, 2008). The bioavailability of these minerals makes sheep milk a valuable source of these elements. Calcium, which is bound to casein in both organic and mineral form, exhibits significant availability during the milk digestion process (Gueguen and Pointillart, 2000); therefore, the calcium bioavailability of sheep milk is strongly associated with high casein levels (Gaucheron, 2005).

Sheep milk's vitamin content is mostly higher than cow's and goat's milk, except for low concentrations of carotene and folate, and pantothenic acid and vitamin D, which have equal concentrations to cow's milk (Park et al., 2007; Wijesinha-Bettoni and Burlingame, 2013). Sheep milk is high in riboflavin, while cow, goat and buffalo milk are adequate sources of riboflavin. As a reference, nutrient intakes of riboflavin (0.5 mg/d) can be achieved with 2 cups of sheep's milk (300 mL). Sheep's milk can also be considered a source of vitamin C, containing an average of 4.6 mg/100 g (Wijesinha-Bettoni and Barbara Burlingame 2013). Since it is not common to drink sheep's milk, likely, two glasses of sheep's milk yoghurt or the equivalent of milk in 90 g of sheep cheese will meet the daily requirements (Recio et al., 2009). Sheep's milk is also a good dietary source of vitamin A and vitamin E. Vitamin A is only found in sheep's milk as retinol because dietary β -carotene is completely converted to this form (Raynal-Ljutovac et al., 2008); However, vitamin E exists in 3 forms, with α -tocopherol being the most abundant form (α -, β -, and γ -tocopherols) (Revilla et al., 2017).

Goat Milk

In terms of composition, it has values close to cow's milk. Since approximately 75% of the proteinaceous substances in its composition are made up of casein, it is included in the group of casein milk. Since the amount of carotene is low, goat's milk is whiter than cow's milk and this is how it is distinguished from cow's milk. The dry matter of goat milk varies between 13-14%. While the amount of non-protein nitrogenous substances is 0.19% in cow's milk and 0.12% in women's milk, it is as high as 0.44% in goat's milk (Uysal and Kılıç, 2005).

The ratio of capric, caprylic and capric fatty acids is high in goat milk fat. Goat milk is 2-3 times richer in vitamin A than other milk. The reason for this is that goats eat more green fodder in winter and the thyroid glands, which play a role in the conversion of carotene to vitamin A, are larger and more active in goats (Veral, 2005).

The lactation period of dairy goats varies between 6 and 10 months. The annual milk yield of culture breeds is around 1,000 litres. This amount increases up to 1,500 – 2,000 litres in specially fed goats. Goats are considered to be the animals that give the most milk in proportion to their body weight, and culture-breed goats give milk approximately 10 times their body weight in a

lactation period. As the yield increases, there is a decrease in the milk-fat and protein ratio (Dellal, 2005).

Goat milk contains a large amount of phosphate. Goat milk is a good source for eliminating phosphate deficiency seen in people who do not have the habit of eating meat and fish. Because it keeps stomach acidity under control, it is recommended that people with stomach problems drink goat's milk. Goat milk is poor in terms of some vitamins, especially vitamin B12, manganese and iron. For this reason, anaemia can be seen in those who are fed goat milk for a long time (Haenlein, 2004).

Breast Milk

The mineral content of breast milk is low and is compatible with the kidney functions of immature newborns. In addition to amino acids, glucose, fatty acids and vitamins, minerals are also needed for the reproduction, regeneration and growth of organisms. Milk protein is of good quality and its utilization rate in the body is 90%. The known contribution of milk proteins to growth and development in the body, as well as their effectiveness in tissue differentiation; It is known that it has positive effects on calcium absorption and immune system functions, reduces blood pressure and cancer risk, is effective in controlling body weight, and is protective against dental caries (Balci, 2011). Lactose, a milk carbohydrate, is the source of milk energy. Lactose is helpful in the formation of brain and nerve cells and in regulating bowel movements. It develops beneficial gut bacteria by providing a suitable environment (pH). Mild nausea, gas in the stomach, heartburn and mild diarrhoea can be seen in those who do not have the habit of drinking milk. These findings go away as you continue to drink milk (Balci, 2011).

Milk fat is another source of milk energy. It provides absorption of fat-soluble vitamins (A, D, E, K). In special cases and adulthood, semi-skimmed or skimmed milk can also be consumed. It has been reported that fatty acids in milk fat are a variable that enables the development of the nervous system and intellectual capacities of children (Samur, 2012).

Milk is rich in minerals (calcium, phosphorus, iodine, sodium, and magnesium). Milk contains more and more available calcium minerals than any other food. Calcium, phosphorus and magnesium are essential components of bone tissue. From childhood to the age of 20-25, with a balanced diet, bone mineral tissue increases. In old age, bone mineral tissue decreases due to inactivity and changes in hormonal balance. Milk, which is rich in calcium, phosphorus and protein content, provides the development of bone tissue in childhood and youth and reduces loss in old age. Milk protein not only increases calcium absorption but also ensures the formation of bone tissue cells. Lactose, a milk carbohydrate, is also an important factor that increases calcium absorption from the small intestines (Coşkun, 2003).

Milk contains a large number of vitamins that have a function in growth and development, the proper use of nutrients in the body, the fulfilment of the functions of the nervous system, and the development of body resistance and blood production. Riboflavin (vitamin B2), B12, vitamin A, B6, B1, niacin and folic acid are vitamins found in sufficient amounts in milk. The vitamins in the composition of milk are extremely sensitive to many physical and chemical effects such as heat and light. Loss of nutrients due to exposure to ultraviolet rays during the processing of milk, especially during heat treatment and transportation, is an undesirable situation in terms of health (Baysal, 2007; Samur, 2012). It is a natural food that is easy to digest and contains fluid, nutrients and energy elements that the newborn needs for ideal development and growth, with high biological benefits. Breast milk in terms of baby and mother; It has many benefits in terms of nutrition, health, immunity, development, psychological and socioeconomic aspects (Hancıoğlu Aytac and Yazıcı, 2020).

The presence of cells, enzymes, hormones, lysozymes, different growth factors, lactoferrin, peptides and special immunoglobulins in the content of breast milk and the high level of taurine

make breast milk a unique compound. In addition to these substances in breast milk, the amounts of iron, zinc, copper, calcium and phosphorus are also adjusted according to whether the baby is premature or not. When the amount of these is adjusted at the beginning, end and different meals and the secretion of special immunoglobulins according to the needs of the baby is taken into account, there is no other explanatory source other than believing that this is made especially for babies by nature (Bilir, 1984; Erbersdobler et al.1984).

CONCLUSION

The amount of milk recommended being consumed for adequate and balanced nutrition of healthy individuals varies according to age, gender and physiological status (growth and development period, pregnancy, lactation, old age). 2-4 (400-800 ml) servings in the Nutrition Guide published by the National Milk and Dairy Products Council and 2 servings for adult individuals in the Turkey-Specific Dietary Guidelines [one serving: one medium glass (200 ml)] It is recommended that children, adolescents, pregnant and lactating women and postmenopausal women consume 3-4 (600-800 ml) portions. Milk contains a large number of vitamins that have a function in growth and development, the proper use of nutrients in the body, the fulfilment of the functions of the nervous system, and the development of body resistance and blood production. When milk and dairy products are considered a source of some important minerals, especially calcium and phosphorus, and some B group vitamins such as protein and riboflavin, it is understood that they are an important food group in terms of public health.

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Immunosensors: Immobilization Methods, Transduction Mechanisms And Current Studies

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Introduction

Biosensors are analytical devices that transfer and interpret signals generated by the immobilization of biorecognition elements (enzyme, hormone, nucleic acid, cell, antibody) on a transducer surface. Biosensors consist of an analyte that is wanted to identify, a biorecognition element or a bioreceptor, and a transducer that convert the reaction which depends on the analyte's specific concentration to a signal (Sharma, 2016). The biocomponents that sense the analytes may be enzymes, antibodies, DNA probes, microorganisms, cells, and tissues while electrochemical, optical, piezoelectric, thermal or magnetic systems are used as transducers. Electrochemical, optical, piezoelectric and thermometric transducers are the four main types of transducers that are used in biosensors. Electrochemical devices usually measure a fixed voltage in a current (amperometry), a voltage in zero current (potentiometry), conductivity or impedance. Impedance can be defined as the resistance to current in an electrical circuit (Tothill, 2001). Optic transducers use some effects like light absorption, fluorescence or refractive index (Tothill & Turner, 2001). Thermometric biosensors work by using enthalpy changes during biochemical reactions. Sensors working with the piezoelectric principle measure the resonant frequency of wave propagation along the piezoelectric material. These principles are used to evaluate mass, viscosity and density changes at the sensor surface (Tothill, 2001). Immunosensors can operate in two ways, according to direct measurement methods, where the detection mechanism is directly dependent on the antigen-antibody interaction, or indirect measurement methods, where a label such as an enzyme or fluorescent molecule is used to detect the binding reactions (Karunakaran, Pandiaraj & Santharaman, 2015). According to the operation of the detection mechanism within the transduction process, immunosensors may be classified as electrochemical, optical, and piezoelectric immunosensors, or based on the use of labels they can be classified as labeled (indirect) and unlabeled (direct) (Holford, Davis & Higson, 2012).

Immunoassays are quantitative analysis methods that utilize antibodies as immunological agents. The selectivity of antibodies is extremely high and antibodies can bind to their specific targets even if there is a high diversity of different molecules in the sample. Immunoassays consist of a group of immunochemical methods and are frequently used in the field of clinical diagnosis (Karunakaran, Pandiaraj & Santharaman, 2015).

Antibodies are produced as the immune system response of hosts exposed to foreign invaders such as bacteria, viruses and parasites called antigens. Antibodies, also called immunoglobulins, are heavy globular plasma proteins. Antibodies can be classified into five classes: IgG, IgM, IgA, IgE and IgD. IgG is the most abundant antibody in serum and also the most commonly used antibody in immunosensor applications. Antibodies bind to the target antigen with

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high affinity, allowing rapid detection of a wide-range analyte spectrum from pathogens to toxins (Karanakaran, Pandiaraj & Santharaman, 2015).

An antibody can be fixed directly to the surface of an electrode or an optical waveguide in the transducer so that it can provide a detectable signal when binding with the analyte occurs. In this case, the antibodies are referred to as primary or captured antibodies. However, antibodies can be used for labeling when they carry labels such as radioisotopes, enzymes, cells, radiant probes, chemiluminescent probes, metal labels or nanoparticles. Antibodies used in this way are also called secondary antibodies or detection antibodies (Li, 2006). Both primary and secondary antibodies in biosensors are generally used in conjunction with electrochemical, optical and piezoelectric transducers. An immunosensor developed for online measurement of progesterone in bovine milk (Delwiche et al., 2001), an unlabeled quartz crystal microbalance (QCM) immunosensor studied for rapid detection of *E.coli* O157:H7 and *Salmonella typhimurium* in food samples (Su & Li, 2004), and an optical immunosensor for the detection of antibiotic and pesticide residues in water (Tschmelak, Proll & Gauglitz, 2005) can be counted among the samples of immunosensors.

Immobilization Methods of Antibodies

The performance of an immunosensor depends on three critical factors. These factors are: biosensing material immobilization without activity loss; analyte accessibility to biosensing materials; and low nonspecific adsorption with the solid substance (Sharma, Byrne & O’Kennedy, 2016). In order to ensure conformational stability and bioactivity during the immobilization procedure, the main goal is to make the active part accessible to the targets. Proper immobilization should be quick and convenient and should prevent biomolecule leaks. In addition, there are several parameters that affect immobilization such as biomolecule size, shape, working area, polarity, functional group presence, and storage conditions. Chemical and physical properties of transducers are also effective on immobilization (Asal et al., 2019).

The compatibility of antibodies to the sensor surfaces can be controlled by interactions between the antibody and surface reagents. Various approaches aim to make the sensor surface more suitable for immobilization with antibodies of specific orientation, such as functionalizing the surface with specific groups (glutaraldehyde carbodiimide, succinimide esters, malenimide or periodate), or designing it with nanostructures (Kierny, Cunningham & Kay, 2012).

Antibodies can be adsorbed on the sensor surface by noncovalent forces such as electrostatic or ionic bonds, hydrophobic interactions, and van der Waals forces. Adsorption may be classified as physisorption and chemisorption. While physisorption includes van der Waals forces and hydrophobic interactions, chemisorption basically includes the electrostatic interactions between the antibody and the surface (Sharma, Byrne & O’Kennedy, 2016). Antibody trapping on conductive polymer films is another noncovalent approach. The capture of antibodies against human serum albumin on galvanostatically polymerized pyrrole was reported (John et al., 1991).

Polyacetylene, polythiophene, polyaniline, polyindole and polypyrrole are common conductive polymers used in immunosensor manufacturing due to their superior biocompatibility against neutral flowing solutions. Polyquinon is another smart material that is used for the immobilization of biomolecules. In addition to their biocompatibility, they have conjugation and transduction capabilities. Thus, it makes it unnecessary to add a redox label to the solution or to the antibody (Sharma, Byrne & O’Kennedy, 2016).

Chemical immobilization is provided by primary bonds such as covalent bonds. Covalent bonds provide stable and irreversible immobilization and are widely used in antibody immobilization. Most covalent immobilizations use functional groups naturally found in

antibodies. Sugar domains and disulfide bonds between two heavy chains are the most frequently used functional groups to ensure correct orientation (Park, 2019). Covalent bonds are formed by surface modifications that enable the interaction of subsequent antibodies with reagents such as hydroxy, thiol, carboxy and amino groups on the surface. Various surface modification techniques have been developed by chemical modification, photochemical grafting, plasma gas discharge and ionized radiation graft polymerization methods. Binding of antibodies to the sensor surface using amine groups is often performed in the form of covalent bonding with the amino acid lysine to the side chains of antibodies, since these groups are relatively easy to access (Sharma, Byrne & O’Kennedy, 2016).

The primary amine groups of antibodies are covalently bound to the reactive succinimide esters. Feyysa et al. (Feyysa et al., 2013) reported that immobilization of anti CRP antibody via amine covalent bond in a microfluidic platform, enhanced the sequence signal compared to passive binding. The manufacturability of this biochip appears to be similar to that of the human CRP ELISA detection kit. Rahman et al. (Rahman et al., 2007) immobilized a polyclonal antibody onto a carboxylic acid group functionalized on the nanoparticle with the quartz crystal microbalance principle for the detection of bisphenol A.

Antibodies are glycoproteins and contain polysaccharides in their heavy chains. These sugar chains are positioned in the Fc region and immobilization over these regions may be the right choice. Saccharide molecules contain many hydroxyl groups and these groups can be converted to aldehyde groups by oxidation. Therefore, sugar chains that contain active aldehyde groups can be used as an immobilization site. (Park, 2019).

Another common method of immobilization on sugar chains is boronic acid immobilization. Boronic acid forms boronate esters with cis-diols of saccharide molecules at room temperature (Park, 2019). Lin et al. provided antibody conjugation on magnetic nanoparticles with the help of boronic acid by covalent binding of the carbohydrate part of the constant region (Lin et al., 2009).

The method of using disulfide bonds between two heavy chains is also used to increase the accessibility of the active sites of antibodies. The disruption of the disulfide bond occurs by agents such as beta-mercaptoethanol or dithiothreitol. Thus immobilization can be provided by utilizing the disulfide bond (Park, 2019).

Covalent immobilization of antibodies over chemical binders such as glutaraldehyde can damage the antigen binding site and reduce antigen binding affinity (Park, 2019). To overcome this problem, molecules with an affinity for antibodies have been studied. Among these molecules, biotin-avidin/streptavidin, protein A and G are frequently used (Sharma, Byrne & O’Kennedy, 2016; Park, 2019). De Juan Franco et al. developed a new, simple and fast orientation method by utilizing a fusion protein region called protein A gold binding site (PAG). Biotin-labeled antibodies can easily be captured by functionalized surfaces with proteins produced by avidin, streptavidin or a combination of these two proteins and form conjugates (De Juan-Franco et al., 2013). Ionescu et al. developed a highly sensitive direct impedimetric immunosensor for the detection of the herbicide atrazine. Affinity-based immobilization of anti-atrazine antibodies on polypyrrole film was performed using a gold electrode with nitrilotriacetic acid. They were able to detect the charge transfer resistance even with atrazine in the amount of 10 pg/ml, and they were able to gain functionality in the range of 10 pg/ml-1 µg/ml (Ionescu et al., 2010).

One of the most promising methods is the molecularly imprinted polymer method. Molecular imprint-based receptors are designed to selectively identify target molecules, including proteins. Molecularly imprinted polymers are generally formed around the template molecule by a

matrix polymerization method. After polymerization, the template is extracted and a porous structure is produced with the dimensions and shape of the molecule to be immobilized, providing an ordered orientation and recognition with high specificity (Kalecki et al., 2020).

Transduction Mechanisms of Immunosensors

Electrochemical

Antigen-antibody interactions occurring in electrochemical transducers cause changes in ion concentration, potential, current, conductivity, capacitance and impedance measurements. Electrochemical biosensors that take advantage of these changes have many advantages, such as being economical, easy to apply, portable, reliable, fast, powerful, mass-produced and functional with small amounts of analyte. Electrochemical immunosensors have become interesting and sought-after biosensing devices with the use of modern and developing technology (Hayat, Catanante & Marty, 2014).

Changes in the dielectric properties of the electrode surfaces can be monitored by potentiostatic procedures. With this discovery, amperometric biosensors have become more popular. Amperometry measures both the magnitude and intensity of the current at a fixed potential value at a given time in an electrochemical cell. When a regulated potential is applied between the electrodes, redox reactions occur that produce a measurable current proportional to the concentration of the relevant analyte (Hayat, Catanante & Marty, 2014). The electrodes used in amperometric sensors have the property of exhibiting high stability over long time periods and showing a linear response in the physiological range. Resulting from molecules such as potassium ferrocyanide being used to mediate the efficient transfer of electrons between the electrode surface and the labels, the electrodes are not affected by the oxygen flow, dissolved gas effects are avoided, and thus stable and accurate results are obtained (Grieshaber et al., 2008). Pinacho et al. developed an enzyme-labeled amperometric immunosensor immobilized on a magnetic bead that detects 7 different fluoroquinolones in milk (Pinacho et al., 2014). When the reaction takes place with the fluoroquinolone samples, the magnetic beads are easily captured by the magnetic graphite-epoxy composition (m-GEC) electrodes. Magnetic beads work specifically for fluoroquinolones from the multi-component structure of milk, hence there is no need for an extra extraction process. Voltammetry is the process of measuring the corresponding change in current while the potential is changed over a predetermined range. As a result, the measured current is translated into peaks and plateaus that represent the target of interest and where the height of the peak gives the amount of analyte. Detection of different multiple analytes in a single sample is possible because each analyte has a characteristic peak position, making the voltammetry technique extremely valuable for immunosensors (Su et al., 2011). This method is extremely sensitive due to low background noise generation, especially when applied after sample preconcentration (Velusamy et al., 2010). Cyclic voltammetry is an important technique that works with the mechanism of charge transfer reactions of redox labels, enabling the determination of biomolecule concentrations depending on oxidation/reduction pathways. In cyclic voltammetry, the potential of an electrode is drawn from the initial potential to the final potential and back to the initial potential. The changing current is measured during these potential transitions (Karunakaran, Pandiaraj & Santharaman, 2015). Singh and Krishnan developed a voltammetric immunosensor that can measure serum insulin level with 5 pM limit of detection. By exposing the analyte to magnetite nanoparticles, they ensured high sensitivity capture of serum insulin conjugates by the electrodes, thus developing the first voltammetric diabetes sensor (Singh & Krishnan, 2015).

Potentiometric measurement records and displays ion activities in an electrochemical reaction. It indicates the mass charge potential of an indicator electrode versus a reference electrode where the current is zero. The Nernst equation can be used to determine the relationship between

the potential and the amount of analyte. The most commonly used components in potentiometry are pH electrodes. Other electrodes like ion (F^- , I^- , CN^- , Na^+ , K^+ , Ca^{2+} , NH_4^+) or gas (CO_2 , NH_3) selective electrodes may also be used for potential measurements (Bobacka, Ivaska & Lewenstam, 2008). The antibodies and antigens in solution each show a net polarity of electrical charge that correlates with the ionic composition of the solution and the molecules it contains. If the antibody complex combines with the antigen, a different electrical charge arises from the charge that the antibody alone has. This change can be measure by potentiometric methods (Karunakaran, Pandiaraj & Santharaman, 2015). Tarasov et al. developed a potentiometric immunosensor that detects BHV-1 specific antibody in bovine serum with a microfluidic chip immobilized with Bovine Herpes Virus-1 viral protein gE. This method, which takes less than 10 minutes to give results, has similar detection limits with ELISA and it is advantageous because it gives fast results (Tarasov et al., 2016).

Conductometric immunosensors measure the changes in conductivity of sample solutions due to biorecognition reactions. Most reactions produce ionic species concentration changes that cause electrical conductivity or current changes. A conductometric immunosensor consists of two electrodes placed at a certain distance from each other and an alternating voltage is applied which causes a current to be generated across the electrodes. If the immunosensor has been exposed to the analyte, the change due to the receptor-analyte combination in the identification layer causes a significant change in the total capacitance (Karunakaran, Pandiaraj & Santharaman, 2015). Mutlaq et al. developed a conductometric-based immunosensor for the detection of *E. coli* O157:H7 bacteria, one of the most common foodborne pathogens. They coated the gold electrode with polyaniline/zinc oxide (PANI/ZnO) nanocomponents, immobilized monoclonal anti-O157:H7 antibody as a biorecognition element, and measured in the detection range of 101-104 CFU/ml in 30 minutes (Mutlaq et al., 2021).

Impedance is basically the opposite of current in an electrical circuit. The difference between resistance and impedance is that the resistance obeys Ohm's law and opposes the current of electrons in alternating and direct current circuits. Impedance is a resistance that changes according to inductive and capacitive effects seen only in alternating current circuits (Leva-Bueno, Peyman & Millner, 2020). In electrochemical impedance spectroscopy (EIS), changes in current are measured after the application of a voltage with a specific frequency to the interface. The current/voltage ratio gives the impedance (Velusamy et al., 2010). C-reactive protein (CRP) is a routinely used biomarker in the clinical diagnosis of liver or cardiovascular inflammation. The amount of CRP in the blood plasma can also be used in the diagnosis of diseases such as diabetes and cancer (Lim & Ahmed, 2019). Jampasa et al. produced an electrochemical biosensor for the detection of CRP with a sandwich model, using a serigraphic graphene electrode modified with cysteine and gold. Anthraquinone was chosen as a redox label because of its stability, simplicity and compatibility with biomolecules. The produced immunosensor measured CRP in the range of 0.01-150 $\mu g/ml$ with 1.5 ng/ml limit of detection (Jamasa et al., 2018).

The sensitivity of immunosensors can be increased by the modification of the surfaces of electrodes by nanomaterials. Because electrode surface area of would increased (Ahmed et al., 2014). Li et al. used gold nanoparticles (AuNP) to detect human epidermal growth factor 2 (HER2), a breast cancer biomarker. In this design, AuNPs are used as support surfaces for immobilization of HER2 specific peptide sequences and since they form a wide surface area they produce a powerful electrochemical current. The group reported 0,5 pg/ml limit of detection and 1 pg/ml to 1 ng/ml detection range (Li et al., 2018).

Optical

Optical immunosensors utilize methods such as chemiluminescence, light absorbance, fluorescence, phosphorescence, light polarization, and rotation for the detection of analyte. Optical sensors work on the principle of detecting changes in light reflections and emission caused by the formation of the Ab/Ag complex, using light as a stimulus. A typical optical sensor receives the light from the laser, diode or white bulb and allows the changes to be observed while reflecting it from the sensor. Measurements can be made sequentially or simultaneously by sending light to the sensor at various angles. Many publications also focus on the use of nanoparticles to improve the performance of optical immunosensors. Nanoparticles generally lower the detection limit and increase signal amplification depending on the increased surface area (Karunakaran, Pandiaraj & Santharaman, 2015).

Optical change can be created with transducer types such as surface plasmon resonance (SPR), interferometry, reflectometric interference spectroscopy, ellipsometry and total internal reflection fluorescence (TIRF) (Abdolrahim et al., 2015).

Optical immunosensors are basically classified into two groups as direct and indirect. A direct optical immunosensor consists of a physical component, a biological component and a detector. While the prism and light source are physical components, the recognition element or biochemical content is biological. These systems generally include an optical reader, a biorecognition material, and sample processing systems, as well as four subsystems: light source, detector module, data processing unit, and optical connections. In the biological component, biomolecules are used as molecular probes for bioconjugation with the analyte. In the physical component, excitation light creates a resonance in the surface plasmons in the film and the reflection intensity of the light reflected at certain angles is observed. The electronic signal which is converted from the reflected light by a linear array detector makes possible monitoring the biochemical interaction between the analyte and the transduction surface. Such tests are completed in about ten minutes (Lee et al., 2018).

A surface plasmon is an excitable electron oscillation that occurs between two surfaces whose insulating constants are of opposite signs. In a surface plasmon resonance (SPR) immunosensor, antibodies are immobilized on a thin metal sheet, usually gold, and detection is made by observing the reflection angle of polarized light with a prism depending on the formation of the target ligand. The metal film acts as a mirror, reflecting this light, increasing the power of reflection and making it calculable and measurable. When immobilized antibodies bind to targets, a deviation in SPR angle is observed depending on the concentration of the target (Lim & Ahmed, 2019). Battaglia et al. developed two SPR-based immunosensors for the detection of procalcitonin (PCT), which is used as a sepsis marker in both human and veterinary medicine, for use in canines and equidae. They utilized the molecular imprinting method using dopamine and norepinephrine to create PCT binding sites on the gold chip surface. In the detection of markers in plasma, these two biosensors developed for canine and equidae reached the of 30 ng/ml and 15 ng/ml limit of detection, respectively (Battaglia et al., 2021).

In indirect methods, fluorophores, luminophores, chromophores, quantum dots and nanoparticles can be used as labels. Total internal reflection fluorescence (TIRF) microscopy can be used to detect excited fluorophores within a 200 nm long field. A laser beam with an angle of incidence greater than the critical angle travels through the silica layer and is reflected from the glass coating towards the inside. Due to the splitting between the transmitted and reflected rays in this way, a standing wave is produced from the silica layer, resulting in an area where waves with decreasing amplitude are seen (Lee et al., 2018). Käppel et al. developed a TIRF-based immunosensor that measures progesterone from milk samples taken from oestrus-cycled cows.

They covalently immobilized the monoclonal antiprogesterone antibody to the sensor surface as a biological identification element and aimed to detect progesterone. As a result, they were able to measure at the picogram level in less than 5 minutes, thus making it possible to monitor critical parameters such as estrus and pregnancy follow-up, which are important for milk production (Käppel, Pröll & Gauglitz, 2007).

Optical fiber immunosensors are another method that allows monitoring of antigen-antibody complexes immobilized on an optical surface. Optical fiber immunosensors do not require reference electrodes, are convenient, and are widely used in clinical diagnosis, in vivo observations, and the detection of hazardous materials. The total internal reflection principle is the basis of optical fiber biosensors. The rays are transferred along the center of the fiber with negligible light loss (Lee et al., 2018). Immunochemical agents are immobilized at one end of the optical fiber and measured by light absorbance or fluorescent emission. In other words, antigen-antibody binding causes a colorimetric change. In this way, measurements can be made depending on the amount of antigen present (Karunakaran, Pandiaraj & Santharaman, 2015). Yang et al. developed a dual channel optical immunosensor that measures SARS-Cov-2 and Influenza A viruses in 10 minutes on a single sensor with a dual channel system. This sensor can recognize nucleoproteins of influenza A H1N1, H3N2 and H7N9 and SARS-CoV-2 Omicron and Delta variants (Yang et al., 2022).

One of the most important nanostructural detection tools is quantum dots, which are nanoparticles consisting of fluorescently labeled probe biomolecules. Quantum dots are semiconductor nano-sized (2-10 nm) crystalline structures. They are brighter and more photostable than conventional fluorophores due to their high quantum efficiencies. In addition, their color is directly related to their size. The larger size emits a larger wavelength, giving a color close to red, while those with a smaller diameter give a color close to blue. Therefore, broad absorption spectra and narrow and symmetrical emission scales can be combined and excited by a single wavelength and functional light source to exhibit a multicolored character in disease mapping and detection (Singh et al., 2018). Kerman et al. developed a quantum dot-based sandwich biosensor for prostate-specific antigen (PSA) detection. They immobilized the primary antibodies with protein A on the carbon surface and ensured that the target antigen was captured by these antibodies. After adding secondary biotinylated antibodies to the assembly, a sandwich type immunoassay was created. Then, they added streptavidin-conjugated quantum dots to the system, enabling them to bind with secondary antibodies and detect the presence of PSA due to the presence of emission of quantum dots (Kerman et al., 2007).

In optical detection, functionalized metal nanoparticles with relatively higher stability and lower background noise can be used as an alternative to fluorescent labeling of antibodies. Gold nanoparticles are the most studied and frequently used particles among these metals. It has easy fabrication, high biocompatibility, conductivity, compatibility with surface modification, and optoelectronic properties based on size and shape, which are useful in biosensor development (Li et al., 2010). Mradula et al. used gold nanoparticles in the biosensors they developed for the detection of thyroxine hormone. They immobilized anti-thyroxine antibodies on AuNPs that they functionalized with cysteamine and were able to measure thyroxine with this bioconjugate with a 9.11 pg/ml limit of detection (Mradula et al., 2020).

Piezoelectric

Piezoelectric immunosensors work with the principle of measuring the mass sensitivity of piezoelectric quartz crystals, which varies depending on the antigen-antibody complexes formed. These types of immunosensors are popularly preferred in many sectors due to their simplicity, high sensitivity, specificity, stability, speed and safety (Lim & Ahmed, 2019). The mass change is

detected by quartz crystals, the main component of these sensors. Piezoelectric crystal oscillates at a specific frequency, if there is coupling, when an electrical signal is used at a certain frequency (Terry, White & Tiqwell, 2005). When voltage is applied to the quartz crystal by two electrodes, the orientation of the crystal changes and mechanical oscillation is observed at a characteristic vibrational frequency. With the formation of immunocomplexes, the crystal's mass increases because of the immunocomplexes extra mass load, the oscillation frequency of the crystal changes, and in this way, mass change can be detected electrically. Measurement methods used in immunosensors can be direct or indirect. In direct measurement of the analyte, the mechanical oscillation frequency decreases with increasing analyte concentration. In indirect measurements, the analyte first interacts with the free elements in the solution. This causes the frequency to change inversely with the amount of analyte (Lim & Ahmed, 2019).

QCM immunosensors are formed from a thin quartz plate with an electrode attached to its edges and an antigen or antibody immobilized on its surface. As piezoelectric QCM works by creating an electric field between two electrodes and oscillating the quartz crystal (Kurosawa et al., 2006). The applied electric field creates an acoustic wave that comes perpendicular to the crystal surface and passes through it. These waves provide the necessary impedance at the point where the width of the device reaches several half wavelengths. As a result, the quartz disc undergoes mechanical resonance where the resonant frequency occurs in the space between the two electrodes, depending on the thickness of the quartz disc and the natural velocity of the acoustic wave (Aberl, Kösslinger & Wolf, 1998). Immunological detection of the target analyte causes an increase in mass, which in turn causes a decrease in the resonance frequency in the oscillation. The change in mass can be determined by calculating the difference between the frequency measurement made before the analyte is applied and the measurement to be made after conjugation (Terry, White & Tiqwell, 2005). Kim et al. developed an immunosensor for the detection of canine parvovirus by immobilizing a monoclonal anti-CPV IgG antibody on a gold-plated quartz crystal surface over thiol groups. They successfully detected CPV after stool sample loading (Kim et al., 2015).

SAW immunosensor setups generally consist of a piezoelectric material placed on a silicon substrate and metal electrodes combined with this material to generate mechanical acoustic waves and make them visible in case of a reaction response (Lim & Ahmed, 2019). This type of sensor is based on the production of a specific acoustic vibration with concentrated acoustic energy around the outer surface of a solid material that provides high sensitivity to surface adsorption. The electrical signal is confined by an independent layer to ensure the concentration of acoustic energy and transforms into polarized lateral waves parallel to the surface of the piezoelectric crystal. Since any material in contact with the surface will significantly change the speed and amplitude of the waves, the transducer can precisely detect all specific biological interactions on the surface of the piezoelectric material. When there is an increase in mass as a result of the formation of immunocomplexes, the excited waves traveling along the crystal surface are subject to frequency changes in the motion path. Changes in the resonance frequency proportional to the mass change on the transducer surface can be measured in this way (Fogel, Limson & Seshia, 2016). Toma et al. developed a SAW immunosensor supplemented with gold nanoparticles and used it for the detection of house dust mites. Gold nanoparticles conjugated with streptavidin affinity bind to the biotinylated detection antibody forming a sandwich format with the capture antibody and antigen immobilized on the surface. With the measurements made in this way, the detection limit of 2.5 ng/ml was reached (Toma et al., 2017).

Thermometric

Heat emission or absorption is often observed when biochemical reactions take place. Based on this phenomenon, thermal immunosensors have been developed based on the concept

of measuring thermal change with the formation of a specific antigen-antibody complex (Ramanathan & Danielsson, 2001). All temperature changes caused by the heat released or absorbed can be measured by measuring the molar enthalpy and by summing the energy of the products formed as a result of biochemical reactions. Heat-based detection methods are generally used in combination with enzymes with thermistor properties, since all reactions with the enzyme are accompanied by enthalpy changes (Lim & Ahmed, 2019). It is common practice to combine the flow injection assay (FIA) method, which pioneered the development of thermometric sensors, and the detection of thermal change in enzymatic reactions. Xu et al. developed an FIA-based thermometric ELISA (TELISA) for the detection and measurement of diazepam (DZP). DZP is used by mixing with beverages in various criminal cases, such as theft and sexual abuse. In this study, they produced a stable temperature change signal that was correlated with temperature changes as a result of enzymatic catalysis. The TELISA design is based on the competitive format determination of the thermal response generated by the catalytic degradation of analyte-enzyme conjugates by the respective products. With this principle, the detection limit of 33.71 ng/ml was reached (Xu et al., 2017).

Conclusions

Immunosensors have become popular tools since they meet the need for a sensitive, specific, fast, miniaturized size and inexpensive diagnostic device that can detect a variety of analytes. Significant progress has been made in this area by developing new protocols using up-to-date technology and modern tools. It has been observed that proper orientation and antigen-binding activity increase significantly with the development of recombinant antibodies and the application of appropriate immobilization techniques. The major challenge in the development of immuno sensors goes beyond the laboratory environment to end-user applications. The prevalence of these sensors will increase with the advances in nanomaterials, protocols, versatile designs, powerful sensors and signal generation.

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Glycosylation and Inhibition of Proteins

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Introduction

The term **glycosylation** refers to the incorporation of various carbohydrates into the structures of biologically important molecules such as lipids and nucleic acids, especially proteins. Incorporation into this structure can occur in two different ways, enzymatic and non-enzymatic. If the binding of carbohydrates to proteins occurs in the catalysis of an enzyme and by glycosidic bond, it is known as **enzymatic glycosylation (Glycolization)**. However, if this binding occurs as a result of the reaction of reducing sugars or metabolic derivatives with proteins without enzyme catalysis, it is called **non-enzymatic glycosylation (Glycation)** (Kılınç, 2011; Kayılı, 2016; Aydınhan, 2018). This review will focus more on glycation events and products.

Enzymatic glycosylation reactions (glycosylation)

It is known that in order for proteins to become functional, they must undergo post-translational modifications, including permanent or temporary enzymatic processes, during or after the translation process. One of the most important post-translational modifications that occur frequently during the synthesis of proteins is protein glycosylation. It is known that after translation of many proteins, especially membrane proteins and a significant part of the proteins secreted out of the cell, various carbohydrate units are attached to their structures and these modifications are important for the structure and functions of these proteins. It has also been demonstrated by studies that glycosylation directly or indirectly affects the function of proteins in various ways and gives glycoproteins various functions in vital processes such as cell stimulation, cell-cell interaction and signal transmission (Kılınç, 2011; Kayılı, 2016).

Structural modification events, which occur by the addition of carbohydrate units to proteins, are usually catalyzed by glycosyl transferase enzymes located in the membranes of the endoplasmic reticulum and Golgi organelles inside the cell. In these highly specific enzymatic glycosylation events, glucose in particular, galactose, mannose, fructose, sialic acids, fucose, N-acetyl mannosamine, N-acetyl glucosamine (GlcNAc), N-acetyl galactosamine (GalNAc) and N-acetyl neuraminic acid (NeuAc) such as monosaccharide units participate in the structure of proteins. During these reactions, carbohydrate units are attached to the side chains of asparagine, serine, threonine or hydroxylysine amino acids in proteins with their anomeric carbons by glycosidic bonds. These bonds can occur in 2 different ways as **O-glycosidic (O-linked)** to the hydroxyl groups in the side chains or **N-glycosidic (N-linked)** glycosylation attached to the amide nitrogen (Kılınç, 2011; Yalçın, 2012; Kayılı, 2016; Aydınhan, 2018).

Since glycosylation is part of the natural functioning of the cell, it takes place in a controlled manner. For this reason, its importance can be better understood when the diseases that occur as a result of abnormal glycosylation of proteins are examined. It has been reported Glycosylation plays a role in various biological events, especially in antigen recognition, infection and immunity, and therefore, the relationship of abnormal glycosylation in proteins with various pathological

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conditions, especially congenital glycosylation disorders (CDG), cancer and immune system diseases (Özaydın & et al., 2012; Yalçın, 2012; Kayılı, 2016).

Non-enzymatic glycosylation reactions (glycation)

Non-enzymatic glycosylation of compounds carrying free carbonyl groups; It refers to the reaction of free amino groups of proteins, nucleic acids and lipids with reducing sugars or their metabolic derivatives without enzyme catalysis. Although this spontaneous reaction is more commonly referred to as protein glycation, it is in today known as the "**Maillard reaction**" (Kılınç, 2011; Toprak, 2014; Demirel & Yıldız, 2018; Yalçın & Rakıcıoğlu, 2022).

It was first reported by Louis Camille Maillard in 1912 that reducing sugars in a solution cause the formation of dark colored products (melanoidins) as a result of the reaction of amino acids. The Maillard reaction, which consists of a series of complex reactions that proceed sequentially and simultaneously, consists of three main stages: early, intermediate and late. The Maillard reaction leads to the formation of products with different chemical properties, such as heterocyclic amines, acrylamide, and 5-hydroxymethylfurfural. In the early stage of the reaction; The carbonyl group of reducing sugars such as glucose, galactose, fructose, mannose, ascorbate, pentoses undergoes condensation with the ϵ -amino group, which is the side chain of lysine amino acids in proteins, with the release of water. In this reaction, which takes place non-enzymatically, the "**Schiff base**" intermediate product, which is an unstable and ketimine intermediate, first appears. Although the Schiff base formation that occurs at this stage of the reaction is reversible, it can also undergo an irreversible Amadori regulation, usually after the reaction progresses to the intermediate stage. The proteins formed at this stage are known as Amadori products. In other words, since the formed Schiff base is an unstable product; By adding water back to the structure, the reaction may be reversed or the reaction may spontaneously convert irreversibly to the stable product 1-amino-1-deoxy-2-ketose (ketoamine or fructosamine). The formation reaction of this stable form of ketoamine is called "**Amadori regulation**". When glycation occurs between glucose and lysine residues in proteins, the resulting Amadori product is called fructoselysine. It has been reported that various Amadori products such as fructosamine and fructoselysine, which are formed as a result of the reaction, have circular and flat forms and these forms are in equilibrium with each other in solution. Under *in vivo* conditions, Amadori products have been found to accumulate by irreversibly binding to both short- and long-lived proteins, reaching equilibrium over a period of 15 to 20 days (Figures 1 and 2) (Singh & et al., 2001; Vlassara & Palace, 2002; Kılınç, 2011; Toprak, 2014; Yılmaz & Karabudak, 2016; Aydınhan, 2018; Kerimoğlu & Türk, 2018; Arslan & et al., 2021; Erim, Ergene & Hecer, 2022; Yalçın & Rakıcıoğlu, 2022).

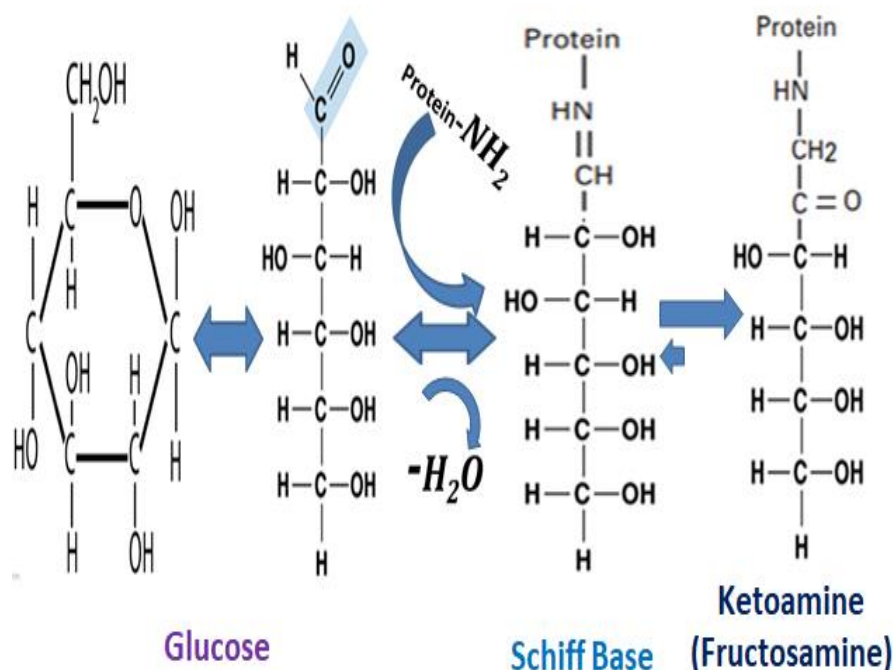


Figure 1. Non-enzymatic reaction of glucose with proteins. The reaction mechanism of other aldehydes and ketones with the amino group is the same. Adapted from Kılınç, 2011.

In the intermediate stage, it is observed that Amadori products undergo oxidation and dehydration reactions and are transformed into various carbonyl derivatives such as glyoxal, methylglyoxal and deoxyglucosons, which are more reactive than the sugars from which they are produced (Figure 2). Studies have shown that dicarbonyl derivatives can arise from the degradation of highly glycated proteins, from glycolysis intermediates, and from peroxidation of lipids (Kılınç, 2011; Yılmaz & Karabudak, 2016; Aydınhan, 2018; Yalçın & Rakıcıoğlu, 2022).

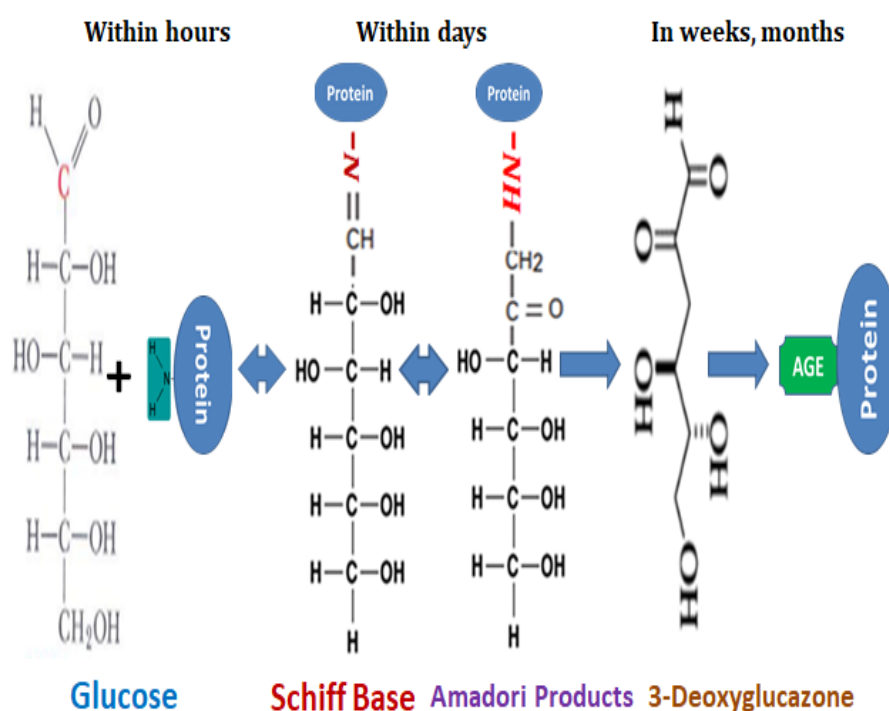


Figure 2. Formation of reactive carbonyl derivatives from Amadori products. Adapted from Toprak, 2014.

In the late stage of the reaction, Amadori products are seen to undergo rearrangements following reactions such as cyclization, oxidation and dehydration to form more stable products. After rearrangements, they form compounds called **Advanced Glycation End Products (AGE's)**, which are generally yellow-brown in color, which can also fluoresce and accumulate in long-lived proteins and cause damage, in an insoluble form with an irreversible reaction (Kılınç, 2011; Aydınhan, 2018; Yalçın & Rakıcıoğlu, 2022).

Although the vast majority of these compounds are known, efforts to understand their chemical properties continue. Recent research shows that Amadori products undergo oxidative and non-oxidative modifications or decompose to form glycation intermediates and AGEs, mainly including furfurals, reductones, dicarbonyls and compounds from the hydroxycarbonyl class (Kılınç, 2011; Aydınhan, 2018; Kerimoğlu & Türk, 2018).

Glycosylated hemoglobin (HbA1c)

Although studies on non-enzymatic glycosylation reactions go back a long time, research on proteins gained momentum with the discovery of hemoglobin forms and elucidation of their structures towards the end of the 1970s. Hemoglobin was the first protein determined to undergo changes as a result of non-enzymatic glycosylation. Studies have focused on HbA, which usually constitutes a large portion (~97%) of total hemoglobin in adults, in which a wide variety of monosaccharide units are bound, such as glucose and fructose. Among the identified forms, HbA1a1, HbA1a2 and HbA1b are respectively attached to the amino groups of the N-terminal valine of the β -chains; They are small amounts of products formed as a result of adding fructose 1,6-bisphosphate, glucose 6-phosphate and unknown carbohydrates to HbA0. One of the most important and most studied forms of HbA1 is HbA1c (Kılınç, 2011; Toprak, 2014; Aydınhan, 2018).

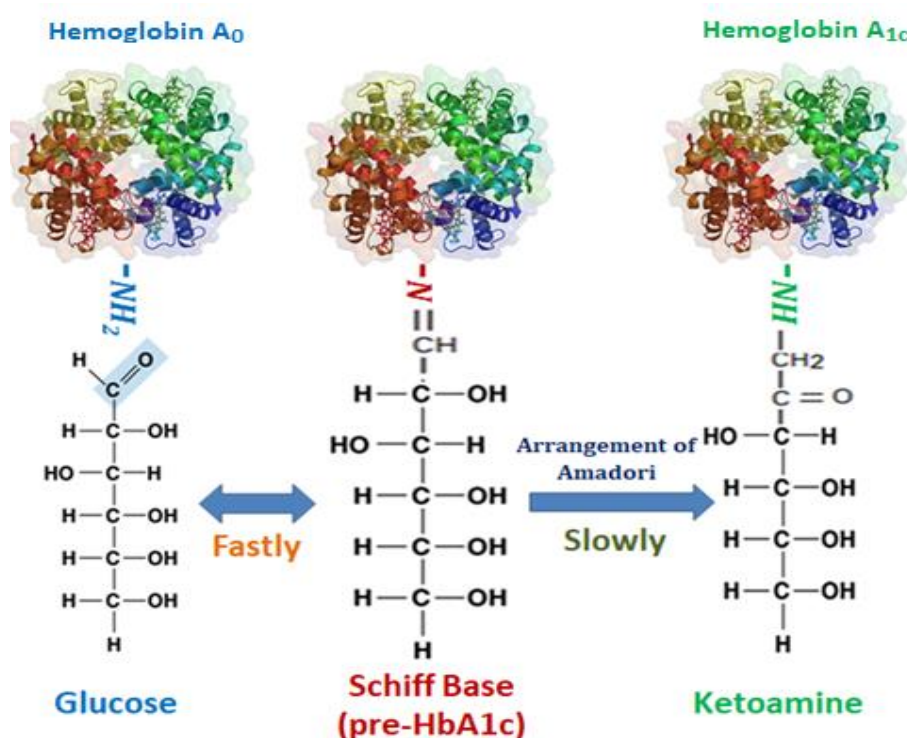


Figure 3. Amadori regulation of hemoglobin and glucose. Adapted from Yılmaz, 2022.

Hemoglobin A1c is the main glycosylated hemoglobin in the blood, accounting for the majority (~80%) of HbA forms. HbA1c is defined as a durable structure formed as a result of non-enzymatic binding of the glucose molecule to the amino group of the N-terminal governor of the β chains of hemoglobin. At the very beginning of HbA1c formation, the structure called pre-

HbA1c occurs as a result of the formation of an aldimine bond (Schiff base) between unstable amino and aldehyde groups. At this stage, the reaction takes place rapidly and bidirectionally; The pre-HbA1c structure is unstable at this time. At this point, the Schiff base can separate or undergo Amadori rearrangement to slowly and irreversibly form the stable ketoamine structure. These stable ketoamines formed by Amadori regulation are called advanced glycation end products (AGEs) (Figure 3) (Kılınç, 2011; Toprak, 2014; Aydınhan, 2018; Yılmaz, 2022).

Glycation of lipids

Since glycation is a reaction between carbonyl groups and amino groups, it was thought for years that only phosphatidylethanolamine (PE) and phosphatidylserine (PS) lipids could participate in glycation reactions. However, contrary to this idea, many *in vivo* and *in vitro* studies have shown that only PE can undergo glycation reactions. As with protein glycation, the level of lipid glycation can also vary depending on the glucose concentration in the medium. Studies have shown that the plasma glycemic PE concentration of diabetics can increase up to 2-3 times compared to individuals with a normal glycemic index. In addition, experimental animal studies have shown that PE glycation occurs in foods that have been processed and stored for a long time. Amadori products, which are formed as a result of glycation of phosphatidylethanolamine, can cause the formation of reactive oxygen species (ROS) by undergoing oxidation and destruction since they are not stable compounds. It has been reported that reactive oxygen species also cause peroxidation in membrane lipids and contribute to the abnormal changes that occur in membrane lipids in various pathologies, especially diabetes and aging (Kılınç, 2011).

Glycation of nucleic acids

Over the years it has been discovered that nucleic acids, like proteins, are non-enzymatically glycosylated. It has been understood that glycation in nucleic acids occurs with reducing sugars over bases containing free amino groups in their structures such as adenine, guanine and cytosine. Glycation reactions have been studied in detail *in vivo* and *in vitro*, focusing more on proteins. Studies involving the glycation of nucleobases have been highly limited to *in vitro* studies, and it has been proven that glycation occurs again with the Amadori reaction. On the other hand, the difference with proteins was observed not in the formation of glycation products, but in the degree of accumulation of the formed products. Accordingly, glycation products formed as a result of glycation of nucleobases show a lower level of accumulation in the cell than products formed by glycation of proteins. The reason for this situation is that the free nucleotides that undergo glycation can be rapidly excreted from the body. In addition, it is thought that the short-lived RNA reduces the glycation rate, and in the longer-lived DNA, the interaction of the bases forming the helix structure through hydrogen bonds is important in protecting nucleobases against glycation. This restriction of glycation of nucleobases has led to the fact that the glycation of these molecules can only be investigated *in vitro*. Studies have shown that guanine bases in DNA are prone to glycation, but they can be separated by hydrolysis due to the instability of the N-glycosidic bond in the formed glycation products. It is thought that the apurinic DNA regions formed as a result of hydrolysis are repaired by the normal DNA repair mechanism and therefore, although the DNA is very long-lived, the accumulation of glycation advanced products on DNA at significant levels cannot occur (Kılınç, 2011).

Factors affecting glycation reactions

Since glycation reactions occur independently of enzymes; The reaction rate is affected by *in vivo* variables such as cellular glucose permeability, the concentration, half-life and reactivity of the reducing sugar and protein to be reacted in terms of free amino groups. Advanced glycation end products can occur endogenously as a part of metabolism, or they can be taken exogenously through food and smoking. It is known that advanced glycation end products occur naturally in

unprocessed raw foods of animal origin, and the cooking process increases the reaction rate by providing more AGEs in foods. It has been reported that cooking methods applied with high temperature for a long time, especially grilling, frying and roasting, increase the formation of AGEs in foods. The rate and amount of AGE formation in foods *in vitro* conditions; It has been proven that many factors such as the level and duration of heat applied to the food, pH in the environment, humidity, composition of the food, food additives and whether the marination process is done or not. In addition to the fact that protein glycation is a slowly progressing reaction, it has been reported that the glycation rates of different amino groups on the same protein molecule may differ from each other. For this reason, since glycation occurs at low levels in proteins with short half-lives, it naturally becomes difficult to detect in these proteins. On the contrary, in proteins with a long half-life, the amount of reducing sugar accumulated on the protein also increases over time, making it detectable. For this reason, proteins with long half-lives such as alkaline phosphatase, elastin, collagen, lysozyme and hemoglobin are generally preferred in *in vivo* protein glycation studies. It has been reported that *in vivo*, protein glycation rate and level vary depending on the half-life of the protein and the concentration of the reducing sugar and free amino group to which the protein is exposed, regardless of the age of the individual (Kılınç, 2011; Yılmaz & Karabudak, 2016; Aydınhan, 2018; Demirel & Yıldırım, 2018; Döner, 2021; Demirel & Yardımcı, 2022; Erim, Ergene & Hecer, 2022).

Advanced glycation end products (AGEs)

Advanced glycation end products are compounds formed as a result of non-enzymatic reactions between reducing sugars and amino groups in the structure of proteins, lipids and nucleic acids. Studies have also revealed different mechanisms of AGE formation. Formation of advanced glycation end products; It has been reported that it occurs by three different mechanisms: Maillard reaction, polyol pathway and increased oxidative stress, and α -dicarbonyl compounds such as glyoxal, glycolaldehyde, glyceraldehyde, methylglyoxal, 3-deoxyglucosone occur in all of them. The polyol pathway is known as a pathway that is activated in the presence of high concentrations of glucose, resulting in fructose. In the polyol pathway, sorbitol, formed by the reduction of glucose by aldose reductase, is converted to fructose by sorbitol dehydrogenase. As a result of overstimulation of the polyol pathway, metabolites such as fructose and triose phosphates accumulate and the intracellular Nicotinamide adenine dinucleotide (NAD⁺) concentration decreases. It has been discovered that the resulting fructose is then converted into products such as fructose-1-phosphate and fructose-6-phosphate, leading to the formation of α -dicarbonyl compounds and contributing to the *in vivo* AGE pool. On the other hand, another mechanism leading to AGE formation is auto-oxidation triggered by an increase in oxidative stress level in the cell. In this mechanism, it has been reported that the formation of α -dicarbonyl compounds occurs as a result of the oxidation of monosaccharides such as glucose, ribose, galactose and fructose by catalytic metals and oxygen. (Döner, 2021; Yalçın & Rakıcıoğlu, 2022; Yılmaz, 2022).

Since Amadori products resulting from glycation are unstable products, they are converted into glycation end products with a large number of glycation intermediates by changing or breaking down by various mechanisms. Glycation can occur by non-oxidative mechanisms, or it can occur with oxidation, and in this case, glycoxidation products are formed. As a result of these mechanisms, it is observed that various compounds, primarily furfurals, reductones, dicarbonyls and hydroxycarbonyl derivative products, are formed. Studies show that these compounds can also cause radical formation by entering redox reactions in the presence of transition elements such as iron and copper. On the other hand, since the reactive products produced by oxidative destruction can react with amino groups in proteins, they form brown macromolecule complexes known as melanoidins by forming covalent cross-links between proteins. In addition, with the oxidative degradation of Amadori products bound to proteins, products such as glyoxal, methylglyoxal, glycolaldehyde, 3-deoxyglucosone, 1-deoxyglucosone, 4,5-dioxopentose and 5,6-dioxohexose can

be formed either free or bound to proteins in the glycation medium (Kılınç, 2011; Toprak, 2014; Muthyalaiiah & et al., 2021).

As 4,5-dioxopentose and 5,6-dioxohexane products formed by the oxidative degradation of Amadori products are involved in covalent cross-linking between proteins, they are generally found bound to proteins; They are the precursors of the two AGE molecules pentosidin and glucospan. The resulting products, glyoxal and methylglyoxal, are imidazole derivatives that cause an increase in cross-linking between proteins in cells and tissues. These compounds formed; It can also occur during the breakdown of amino acids such as threonine and serine, and triosephosphates, especially in the breakdown of Amadori products and lipid peroxidation intermediates. Carbonyls and dicarbonyls, which are among the reactive intermediates formed, cause covalent cross-linking as a result of reaction with lysine/arginine side groups and N-terminal amino groups in proteins. Covalent cross-linking can occur on the same protein or between different proteins. Although it is not fully understood yet, it is thought that the formation of cross-links takes place in a very complex way and lysine residues are of great importance in this formation (Kılınç, 2011; Toprak, 2014; Muthyalaiiah & et al., 2021).

Today, nearly 40 types of AGEs have been defined in free and bound form. Various advanced glycation end products that have been shown to accumulate in tissues have been identified by studies, and as the number of AGEs discovered over time has increased, it has been necessary to classify them. When classifying advanced glycation end products, they are divided into four different groups according to their ability to reflect fluorescent light and their biochemical properties (Döner, 2021; Yalçın & Rakıcıoğlu, 2022):

1. Fluorescent and cross-linking molecules
2. Non-fluorescent and non-crosslinking molecules
3. Non-fluorescent and cross-linking molecules
4. Molecules that fluoresce and do not crosslink

For example; Pentosidine is a fluorescent glycoxidation product formed by cross-linking of arginine and lysine residues with ascorbic acid or monosaccharides. Apart from pentosidine, pentodilysin, AGE-IX, vesperlysin A and C are other AGE molecules known to be cross-linking and fluorescent. The methyl glyoxal-lysine dimer (MOLD) and glyoxal-lysine dimer (GOLD) formations, which are formed as a result of the reaction of methyl glyoxal and glyoxal with lysine amino groups, are cross-linking but non-fluorescent AGE molecules (Figure 4). Although it has been reported that imidazolium cross-links are formed with lysine groups during the formation of these two AGE types, it has been reported that they can also be cross-linked with arginine groups (Two AGEs: GODIC and MODIC). Carboxy methyl lysine (CML), carboxy ethyl lysine (CEL), imidazolone and pyralin are advanced glycation end products that are determined to have no cross-linking and fluorescent properties. Carboxy methyl lysine is widely used in studies to evaluate the level of non-fluorescent AGEs in plasma and bone tissue. Although it can occur as a result of oxidation in various mechanisms, CML can also occur directly as a result of the reaction between the amino groups of glyoxal and lysine. Pyraline, on the other hand, is an important AGE molecule, which has a ring structure and is formed as a result of the reaction between lysine and glucose. The most important AGE molecule, which is not cross-linked but has fluorescent properties, is Argpyrimidine; It has been reported that it may occur as a result of the reaction between methyl glyoxal and arginine, and its serum levels are detectable in individuals with diabetes (Kılınç, 2011; Toprak, 2014; Demirel & Yıldırım, 2018; Döner, 2021; Muthyalaiiah & et al., 2021; Yalçın & Rakıcıoğlu, 2022).

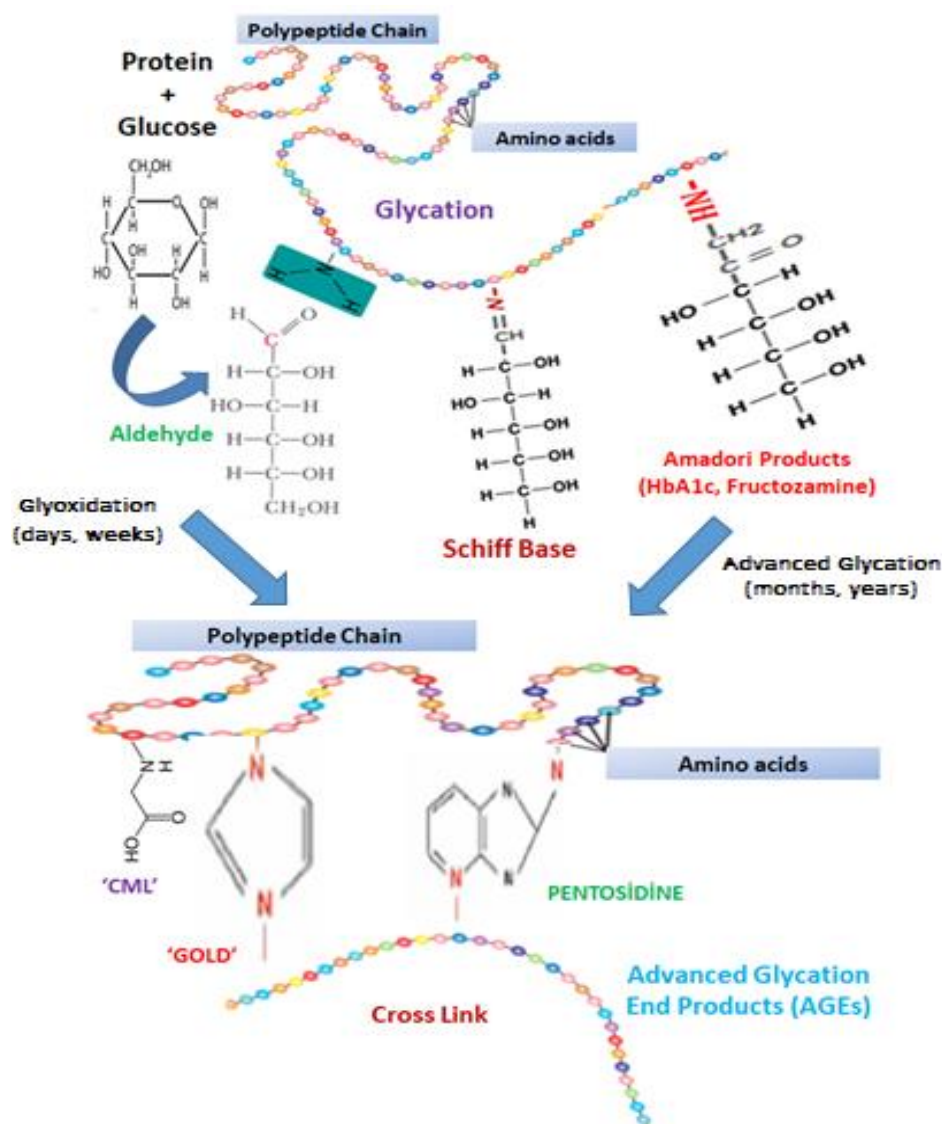


Figure 4. Formation of advanced glycation end products (AGEs). Adapted from Kerimoğlu & Türk, 2018.

RAGE (Receptor for AGEs)a

The undesirable *in vivo* effects of advanced glycation end products are thought to occur in two different ways. It is thought that the first of the exit mechanisms is the change of tissue or vascular structure after the formation of cross-linked structures, and the other mechanism arises through AGE-sensitive receptors. Different receptors with which advanced glycation end products interact in many cell types such as leukocytes, macrophages, neuronal cells and muscle cells have been identified. Among these receptors; **receptors for advanced glycation end products (RAGE)**, scavenger receptors (Class A, CD36, Class B tip 1, LOX-1, FEEL-1, FEEL-2), oligosaccharyl transferase-48 (AGE-R1/OST-48), AGE-R2/80K-H and Galectin-3 (AGE-R3) are reported to be involved. Among these receptors, the most studied is RAGE, a member of the immunoglobulin G superfamily. Advanced glycation end-product receptors (RAGE) are defined as a transmembrane pattern recognition receptor that can be stimulated by many ligands, belongs to the immunoglobulin superfamily, is located on chromosome 6, has a molecular weight of 45 kDa, and acts as a cell adhesion molecule. It was detected and purified for the first time in bovine lung endothelial cells. It is thought that the accumulation of advanced glycation end products in tissues and organs has important effects on the organism, causing glycative stress mediated by RAGE, and binding of RAGE with its ligands generally increases oxidation and inflammation. Advanced

glycation end products, via RAGE; It has been shown that it has a role in important biological events such as autophagy, endoplasmic reticulum stress, mitochondrial dysfunction, hypoxia, epigenetic modification and cancer, and contributes to the development of various pathologies by inducing or inhibiting various intrinsic cell signaling pathways. It is known that important ligands such as β -amyloid, advanced oxidation protein products (AOPPs), calcium-binding S100 proteins and high mobility group box protein 1 (HMGB-1) can also bind to advanced glycation end-product receptors (Singh & et al., 2001; Demirel & Yıldız, 2018; Yılmaz & Karabudak 2018; Döner, 2021; Muthyalaiiah & et al., 2021; Oshima & et al., 2022; Yalçın & Rakıcıoğlu, 2022).

Genome analyzes including other vertebrates such as birds, amphibians, fish and reptiles have proven that RAGE homologs are only found in mammals. From an evolutionary perspective, this information highlights the importance of understanding why mammals need RAGE and what its physiological roles are. Studies show that RAGE has 2 isoforms known as **soluble RAGE (sRAGE)**. These two isoforms of sRAGE; endogenous secretory RAGE (esRAGE) and cleaved RAGE (cRAGE). Endogenous secretory RAGE (esRAGE) form; It occurs as a result of alternative splicing of RAGE mRNA. Fragmented RAGE (cRAGE); It is a non-membrane-bound extracellular form of RAGE as a result of degradation of full-length RAGE by metalloproteases. It is thought that soluble RAGEs (sRAGEs) act as traps by preventing membrane-bound RAGEs from binding with their ligands and removing AGEs from the circulation (Döner, 2021; Muthyalaiiah & et al., 2021; Oshima & et al., 2022).

Glycation inhibitors

With the understanding of the biological effects of advanced glycation end products, different approaches have begun to be developed within the methods of eliminating the harmful effects they create. One of the most studied approaches among these approaches is to reduce or prevent the formation of AGEs. For the inhibition of the production of advanced glycation end products, compounds that reduce the amount of AGE formation by affecting the formation steps were used; For this, Aminoguanidine, which is a nucleophilic hydrazine compound, and pyridoxamine, which can bind to carbonyl compounds during their formation, have been frequently used molecules over the years (Vlassara & Palace, 2002; Toprak, 2014).

Another method that is the subject of the studies is the breaking of the formed AGE cross-links. Agents known as AGE 'bond' breakers used in studies are defined as compounds that break down AGEs and allow them to be removed from the kidney and liver. Phenyl-thiazolium-bromide (PTB), which can break the carbon-carbon bond between dicarbonyls, is a compound in this group. In addition, ALT-711, now known as alegebrum chloride, has enzymatic properties and is reported to be able to break covalent bonds repeatedly, is another important compound in this group (Vlassara & Palace, 2002; Toprak, 2014).

AGE inhibitors and cross-link breakers act on AGE-related pathologies in different ways. Inhibitors are compounds known to reduce the rate of AGE formation but are unable to break newly formed AGE cross-links on long-lived tissue proteins. It is known that crosslink breakers can break AGE crosslinks but cannot prevent new AGE formations. For this reason, it is thought that the most effective approach in pathological conditions due to advanced glycation and its products may be a combined treatment in which inhibitor and bond-breaking agents are used together (Toprak, 2014).

One of the methods that are the subject of research is the RAGE blockade. In this method, sRAGE and RAGE antibodies are used in studies to prevent AGE-RAGE interaction. The main purpose of this method is to bind AGEs in tissues and prevent them from binding to their receptors, thus preventing RAGE-mediated cell activation (Toprak, 2014).

In addition to *in vivo* AGE production, it is known that AGEs are an important exogenous source and contribute to the *in vivo* AGE pool, especially in heat-treated foods. Therefore, another

research topic has been the focus on reducing exogenous AGE exposure. It did not take long to realize that the reducing sugars found in foods, apart from being able to react with proteins and molecules containing amino groups, also form aldehydes and ketones by spontaneous oxidation without reacting with proteins or by breaking down especially during cooking. It has been understood that the aldehydes and ketones formed can form glycation intermediates and glycation end products by reacting with proteins. Understanding that they are formed in high amounts especially in foods cooked/fried at high temperatures in the studies carried out, has led to a rapid increase in research on this subject. It has been reported that approximately 10-30% of dietary AGEs are absorbed through the gastrointestinal tract after oral intake and enter the circulation. It is stated that approximately 1/3 of the AGEs that enter the circulation are excreted in the urine within 48 hours, and the remainder accumulates in the body.

In studies, the effects of making changes in eating habits or using some agents that inhibit the absorption of AGEs taken with the diet were investigated. Among these agents, the most studied is AST-120, an oral adsorbent. Studies have shown that regular use of AST-120 has positive effects in individuals with chronic renal failure and slows the progression of the disease. It was also stated that after the use of AST-120 in non-diabetic individuals with chronic renal failure, this compound binds to CML and causes a decrease in AGE serum levels (Toprak, 2014; Demirer & Yardımcı, 2022; Oshima & et al., 2022; Yalçın & Rakıcıoğlu, 2022).

Over the years, compounds with antioxidant activity and/or metal chelation have also been tried in studies and their effects have been investigated. Many *in vitro* studies have shown that AGE formation is inhibited by the use of aspirin, ibuprofen, indomethacin, D-penicillamine, desferoxamine and flavonoids. The reduction of oxidative stress of these molecules has been shown as a possible mechanism. In addition, *in vitro* studies have shown that Diclofenac can also inhibit early glycation products, while piaglitazone, a thiazolidinedione derivative, acts on metal chelation by keeping dicarbonyl compounds and has effects on reducing AGE formation (Toprak, 2014; Yalçın & Rakıcıoğlu, 2022).

In addition, it has been reported that positive results have been obtained with the use of agents such as Aldose reductase inhibitors, angiotensin receptor blockers, metformin, thiamine and benfotiamine in studies over the years (Toprak, 2014).

Although it has been accepted for many years that repair activities occur only on DNA, recent studies have revealed that proteins that have undergone different chemical changes, especially glycation, can be repaired by various mechanisms (Kılınç, 2011).

After the discovery that isoaspartate is formed by the reduction of methionine sulfoxide, which is first produced as a result of the oxidation of methionine in biological systems, by the related enzyme; It did not take long to realize that ketoamines, which are formed by the glycation of proteins, can also be repaired by isomerases, oxidases and kinases. Studies show that these enzymes phosphorylate the carbon-nitrogen bond, thereby destabilizing monosaccharides and separating them from proteins. This mechanism, which has been shown to be able to reverse glycation, has been called "**deglycation**". The first enzyme found to be involved in deglycation was fructosamine-3-kinase (FN3K). In this process, fructosamine becomes unstable by being phosphorylated from carbon 3 by FN3K and is separated from the protein. Studies show that FN3K can mediate not only protein deglycation but also the deglycation of free fructoselysine. After the discovery of fructosamine-3-kinase, the existence of other enzymes that lead to the separation of monosaccharides from proteins, resulting in decreased stability of various amadori products, has also been reported (Kılınç, 2011).

It has been reported that glycation causes various pathologies through direct or indirect changes in biochemical, functional and structural properties of proteins, and inhibition of glycation

has positive effects on these pathological conditions and related complications (Kılınç, 2011; Toprak, 2014).

Today, it has been revealed that the causes of complications in metabolic disorders in which hyperglycemia is seen, especially Diabetes Mellitus, are related to glycation and/or advanced glycation end products. Studies continue to reactivate inactive molecules (proteins, lipids, nucleic acids, etc.) resulting from glycation and/or to eliminate glycation end products.

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Pathophysiological Effects of Advanced Glycation end Products

Pınar COŞKUN¹
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Introduction

Advanced glycation end products (AGEs) can be formed as a result of endogenous or exogenous non-enzymatic glycation, by reactions between the reactive carbonyl group of a reducing sugar and nucleic acids, lipids or proteins. For years it was assumed that AGEs were only more stable structures formed by the non-enzymatically binding of a reducing sugar to lysine residues of proteins. However, in recent years, it has been shown that Methylglyoxal, one of the AGEs, may occur as a result of the enzymatic reaction of glyceraldehyde 3-phosphate with cysteine, lysine and arginine residues of proteins. AGEs that can be produced endogenously or exogenously; It has been reported to have undesirable effects on cellular functions in various ways such as the production of free radicals, oxidation of lipids and nucleic acids, changes in enzyme activities, hydrolysis of lipids and proteins, carbonyl stress and interaction with AGE receptors. Two mechanisms are at the root of the pathological conditions and associated complications caused by advanced glycation end products at tissue and cell level in vivo: The first of these mechanisms is that AGEs cause the deterioration of matrix structure and functions by forming cross-links to the proteins in the structure of the extracellular matrix (ECM). The other mechanism emerges with metabolic changes resulting from the synthesis and release of some proteins and cytokines by activating or inhibiting various signaling pathways via AGE-sensitive receptors (RAGE). It has been reported that these receptors can also induce inflammation by being stimulated by inflammatory cytokines, amyloid- β and other fibrillar proteins besides AGEs (Çetiner & Rakıcıoğlu, 2020; Vıcıl & Ulutaş, 2020; Arslan & et al., 2021; Döner, 2021; Rungratanawanich & et al., 2021; Demirer & Yardımcı, 2022).

It is thought that the pathological effects of advanced glycation end products may be due to their ability to directly trigger oxidative stress and inflammation, as well as providing the generation of reactive oxygen (ROS) or nitrogen (RNS) species. It has been reported that these properties are directly or indirectly related to changes in structural and functional properties in proteins, cellular dysfunction, apoptosis, and ultimately disruption of integrity in tissues and organs. In addition, the binding resulting from the interaction of advanced glycation end products with RAGE receptors; It has been discovered that it is an important factor in the development and progression of diabetes, cancer, neurodegenerative diseases, cardiovascular complications, osteoporosis, kidney and liver dysfunctions, and various diseases related to aging (Rungratanawanich & et al., 2021; Oshima & et al., 2022).

The effect of crosslinking

Considering the normal tissue architecture, it is seen that the cross-linking of the proteins in the extracellular matrix (ECM) with each other is of great importance in ensuring and maintaining tissue integrity. However, the formation of this structure consisting of cross-linked proteins is not sufficient alone, and it is an important condition for this structure to have a certain level of flexibility for its functionality and sustainability. It is known that AGE accumulation in cells and tissues can cause various toxic effects. With the accumulation of AGEs directly in ECM proteins, the formation of new cross-links in addition to the existing cross-links occurs, thus reducing the

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flexibility of the proteins. It has been shown that the decrease in flexibility, which occurs due to the accumulation of advanced glycation end products, especially in ECM proteins such as collagen, elastin and laminin, has a role in various pathologies (Demirel & Yıldız, 2018; Bijnen & et al., 2019; Arslan & et al., 2021; Döner, 2021).

The heart is one of the organs most affected by the decrease in flexibility caused by these reasons. Decreased flexibility has been shown to mediate various heart conditions by causing diastolic dysfunction in the heart. It is known that the sarcoplasmic reticulum Ca^{2+} -ATPase (SERCA) and the ryanodine receptor (RyR) are of critical importance for the cardiac muscle to be adequately contracted and relaxed. It has been clarified by various studies that the glycation of these molecules in the aforementioned system and their cross-linking with AGEs inhibit their functions and may lead to deterioration in diastolic functions by reducing cardiac contractility. (Döner, 2021).

Since the formation of advanced glycation end products is usually due to the high amount of glucose in the blood, intermolecular cross-linking that leads to a decrease in the elasticity of ECM proteins such as collagen also occurs at a high rate in hyperglycemic diseases. It has been reported by various studies that the accumulation of AGEs on proteins prevents the proteolytic degradation of proteins and especially the cross-linked collagen becomes resistant to matrix metalloproteinases and causes disruption of the collagen cycle. It has also been shown that inhibiting proteolytic degradation may cause thickening of the capillary basement membrane, resulting in narrowing of the vessel diameter. It has also been reported that the cross-links between type IV collagen and laminin under the influence of AGEs disrupt the spread and adhesion of endothelial cells to the basement membrane. As a result of glycation of the NC1 chain (non-collagenous), which is expected to be attached to the helix-rich chain in the type IV collagen structure under normal conditions, it is not possible to bind to Type IV collagen. It has also been reported that AGE accumulation can inhibit the binding of collagen and heparan sulfate to vitronectin, an adhesive matrix molecule, and inhibit intramolecular interactions and polymerization by disrupting the cross-shaped structure of laminin. These events may lead to the formation of atherosclerosis by disrupting the ECM structure (Döner, 2021; Selvakumar & et al., 2022).

It is reported that one of the proteins most affected by glycation with the formation of cross-links is LDL. It has been reported that as a result of the formation of intramolecular cross-links on LDL by the effect of AGEs, modified clearance of LDL and its uptake into the cell by receptor-mediated endocytosis have been reported. At this point, it has been determined that AGE-modified LDL becomes more prone to foam cell formation and increases the likelihood of atherosclerosis. It was also understood that the rate of degradation and internalization of glycated LDL decreased and became more open to oxidation. It has also been reported that AGE accumulation on LDL can also reduce nitric oxide (NO) formation by inhibiting the reuptake of L-arginine, which is activated due to blood flow tension in endothelial cells (Döner, 2021).

Glycation reactions are reactions that can cause changes at the molecular level by progressing at different rates in different periods of life. Accordingly, glycation products can be defined as products that are formed under physiological conditions, but whose formation is observed to increase with age. For years, aging has been recognized as a major risk factor for the occurrence of osteoarthritis. However, this idea has begun to be better understood with the examination of the role of AGEs, which are caused by the glycosylation of macromolecules, in aging. It is now accepted that osteoarthritis occurs as a clinical and pathological result of a series of disorders that cause limitation of movement and structural and functional disorders of the synovial joints, which are primarily seen in elderly individuals. At this point, it was also effective to elucidate the product accumulation caused by the low level of AGE inhibition in the aging process and the deterioration in ECM metabolism (Demirel & Yıldız, 2018; Karataş, Şentürk, Efe Arslan & Korkmaz, 2021; Yılmaz & Polat, 2022; Zhou & et al., 2022).

Receptor-mediated effects of advanced glycation products

The most important known mediators of the effects of advanced glycation end products on tissue and cells are AGE-sensitive receptors (RAGE). Some metabolic changes occur as a result of the release of certain proteins and cytokines by activating or inhibiting various signaling pathways through these receptors. In addition, it has been reported that RAGE receptors can also be stimulated by inflammatory cytokines, amyloid- β and other fibrillar proteins other than AGEs. It has also been demonstrated that various cell signaling pathways such as p38, MAPKs, ERK, p21RAS, Pi3K and janus kinase (JAK) can be activated as a result of the interaction of advanced glycation end products with RAGE. Finally, various studies show that AGEs have a role in important biological events such as RAGE-mediated angiogenesis, proliferation, inflammation, fibrosis, and thrombosis (Arslan & et al., 2021; Çetiner & Rakıcıoğlu, 2020; Döner, 2021; Rungratanawanich & et al., 2021).

One of the signaling pathways induced by the binding of RAGE with advanced glycation end products is the NF- κ B (nuclear factor κ B) signaling pathway. The NF- κ B family of transcription factors is known as a signaling pathway that plays a role in cells' response to various stimuli. It is also a signal regulator that is associated with many signaling pathways. It has been understood that advanced glycation end products may play a role in collagen degradation by inducing phosphorylation of the NF- κ B and MAPK families. Therefore, it is thought that controlled inhibition of NF- κ B/MAPK phosphorylation may be an important therapeutic approach in the prevention of age-related chondrocyte inflammation and extracellular matrix degradation. In addition, it has been shown that AGE-RAGE interaction can stimulate the synthesis of cell adhesion molecules such as intercellular adhesion molecule-1 (ICAM-1), E-selectin and vascular cell adhesion molecule (VCAM) by inducing NF- κ B, thus impairing endothelial cell function and increasing vascular permeability. . These molecular adhesives, known as "Cell Adhesion Molecules (CAM)" and providing the coexistence of cell clusters in biological structures; They are protein molecules found on the cell surface and involved in cell-cell and cell-matrix interactions. However, the most important feature of the NF- κ B signaling pathway is its ability to induce inflammatory gene expression. It has been understood that NF- κ B stimulation, which occurs as a result of the interaction of advanced glycation end products and RAGE, stimulates the synthesis of many proinflammatory cytokines and the formation of free oxygen radicals as a result. It was understood that these ROSs increased the expression of monocyte chemoattractant protein 1 (MCP-1) and accelerated the migration of monocytes to the subendothelial region by inducing chemotaxis. Apart from the mentioned mechanism, free oxygen radicals can also occur with the activation of NADPH oxidase directly as a result of the AGE-RAGE interaction. Advanced glycation end products-RAGE interaction can stimulate granulocyte-monocyte colony stimulating factor (GM-CSF) synthesis from macrophages. It is also reported that this interaction can induce the synthesis of tumor necrosis factor α (TNF- α), platelet-derived growth factor (PDGF), IL-1 β and insulin growth factor-1 (IGF-1). One of the signaling pathways indicated to be induced by the interaction of advanced glycation end products and RAGE is the Janus kinase/signal converters and activators of transcription (JAK/STAT) signaling pathway. This pathway is one of the leading pathways used to convert a large number of signals for development and homeostasis in animals from humans to flies. In mammals, the JAK/STAT pathway also serves as the main signaling mechanism for a wide variety of cytokines and growth factors. It has been reported that the JAK-STAT signaling pathway activated as a result of the said AGE-RAGE interaction mediates the proliferation of smooth muscle cells in the damaged vessel wall (Rawlings, 2004; Terekeci, Şahan & Top, 2008; Şensoy & Öznurlu, 2009; Ledoux & Perkins, 2014; Döner, 2021; Zhou & et al., 2022).

Anti-aging treatment studies; Focusing on anti-glycation to reduce morbidity, stimulate healthier aging, longevity and faster recovery, it has received and continues to receive a rapidly increasing interest in recent years. At this point, studies; Targeting of RAGE continues with a focus on developing a preventive and therapeutic strategy against numerous RAGE-related diseases,

including diabetes mellitus and its complications, inflammatory disorders, aging-related diseases, neurodegenerative disorders, cancer (Oshima & et al., 2022).

Effects of advanced glycation end products on tissues and cells

In addition to being exogenously taken from the diet, carbonyl compounds can be formed endogenously in the body as a result of protein glycation and lipid peroxidation; Since they have reactive properties, they can react with proteins and cause unexpected changes in the existing functional, biochemical and structural properties of proteins. These changes that occur with the reaction of carbonyl compounds with proteins; As they may arise independently of AGE receptors, these proteins, whose structure changes with these reactions, also have various effects, especially triggering oxidative stress in cells and tissues by binding to AGE receptors (Kılınç, 2011).

The main pathologies determined to be caused by the effects of glycation on proteins at the cell and tissue level directly or through AGE receptors can be generally classified as follows (Kılınç, 2011):

Increased lipid peroxidation: In case of an increase or a decrease in the inactivation of reactive oxygen derivatives (ROS) in cells and tissues, the balance in the body is disrupted and this is known as oxidative stress. Oxygen free radicals are highly reactive because they are molecules containing one or more unpaired electrons in their outer orbitals. The increase in oxygen radicals resulting from oxidative stress is a mechanism that can be triggered in various ways within the cell. Advanced glycation products can cause oxidative stress by forming pro-oxidant molecules in the body. In particular, diabetes is also defined as a state of increased oxidative stress. AGEs, which may occur as a result of hyperglycemia-induced glycototoxicity, may mediate the formation of superoxide by causing activation of NADPH oxidase in the cell. In this case, increasing reactive oxygen derivatives; It can interact with biomolecules such as proteins, lipids and nucleic acids, causing structural or functional changes in proteins, loss of membrane integrity of cells and genetic-based disorders. It is known that cells have some enzymatic and non-enzymatic antioxidant defense systems that can eliminate these radicals in order to cope with the harmful effects of ROS. One of the most important antioxidant enzymes used by cells to resist oxidative stress is the enzyme superoxide dismutase (SOD). However, it has also been discovered that this enzyme can be inhibited as a result of glycation. It has been understood that the induction of oxidative stress by oxidative destruction of glycation products or various activation of AGE receptors reduces nitric oxide (NO) levels, while inducing endothelin-1 synthesis, leading to lipid peroxidation and increased vasoconstriction at the cell and tissue level. It has also been reported in various studies that some AGE inhibitors have positive effects on reducing oxidative stress (Kılınç, 2011; Demirel & Yıldırım, 2018; Çetiner & Rakıcıoğlu, 2020; Arslan & et al., 2021).

Induction of various cellular activities: Various studies have shown that synthesis and release of many proteins and cytokines occur by activating or inhibiting various signaling pathways via AGE-sensitive receptors (RAGE). It has been reported that many metabolic changes occur as a result of this synthesis and release. It has been revealed by various studies that these receptors can also induce inflammation by stimulating with inflammatory cytokines, amyloid- β and other fibrillar proteins besides AGEs. Stimulation of glycated and glycation products and AGE receptors can trigger the synthesis and secretion of cytokines, primarily TNF- α , IGF-1A, VCAM-1 and IL-1. In parallel with the induction of the synthesis of inflammatory cytokines, chemotaxis of mononuclear cells, gamma-interferon synthesis, T-cell activation and mitogenesis can be observed. In this way, it can be seen that as a result of uncontrolled cellular activities, in addition to thickening of the basement membrane, reorganization may occur in the tissues (Kılınç, 2011; Vıcıl & Ulutaş, 2020; Döner, 2021).

Structural and functional changes of proteins: It is known that AGE accumulation in cells and tissues can cause various toxic effects. The formation of cross-links occurs with the

accumulation of AGEs directly in the proteins, and thus, changes in the structure and functions of the proteins are observed. It is known that glycation can change the net charges and three-dimensional conformations of proteins and lead to the formation of protein aggregates resistant to proteases by cross-linking. The best example that can be given to this subject is the advanced glycation of the crystalline protein in the lens tissue. While the formed glycated crystal causes opacification in the eye and vision problems; It is observed that collagens undergoing rapid glycation in individuals with diabetes age prematurely and lose their elastic properties faster. It has been understood that the cell-matrix interactions are changed as a result of glycation of the proteins in the cell membrane and the extracellular matrix, and the adhesion and spreading properties of the cells are also directly and indirectly affected by this process (Kılınç, 2011; Demirel & Yıldırım, 2018; Bijnen & et al., 2019; Arslan & et al., 2021; Döner, 2021).

Effects on thrombosis and fibrinolysis: Thrombosis can also be defined as an abnormal plug formed by blood elements in the vessel. It appears to occur as a result of the disruption of the delicate balance between coagulation and the fibrinolytic system. It has been reported that glycation, in addition to increasing tissue-related thrombosis factors, also reduces thrombomodulin (TM) synthesis. Thrombomodulin is a glycosylated thrombin receptor with a proteoglycan structure, mostly found in endothelial cells. It is so named because it has a regulatory role on thrombin functions. In addition to being an important factor on the delicate balance between coagulation and fibrinolytic processes, it is also known to play an active role in inflammation, embryogenesis, cancer, diabetes and collagen vascular diseases. It has also been shown that glycation plays a role in stabilizing fibrin by increasing platelet aggregation and the synthesis of plasminogen activator inhibitor-1 (PAI-1), a serine protease inhibitor. As a result, the life span of platelets is shortened and their tendency to form aggregates increases. In parallel, it is seen that the sensitivity of fibrin and fibrinogen to plasmin decreases and the functions of antithrombin III are suppressed. As a result, all these chain events result in the disruption of the delicate balance between coagulation and the fibrinolytic system. Disruption of this delicate balance in the direction of fibrinolysis can lead to bleeding; Increased coagulation can cause thrombus formation (Ermiş, Turgan & Ersöz, 2006; Kılınç, 2011; Celkan & Dikme, 2018).

Advanced glycation end products and glucotoxicity

It is seen that the formation of advanced glycation end products generally occurs due to the high amount of glucose in the blood. If this rise in glucose levels above normal physiological concentrations is prolonged, the amount and function of insulin produced by the beta-cells of the pancreas is affected, and thus beta-cell function is gradually deteriorated. This pathophysiological condition, which is frequently seen in diabetic individuals, is generally known as "beta-cell glucose toxicity". AGEs, which occur as a result of hyperglycemia-induced glycototoxicity and are also defined as glycotoxins; It has a series of chemical effects on cells and tissues, and it has been demonstrated by various studies that it has a role in many pathologies, including diabetes-related complications, atherosclerosis, kidney and neurodegenerative diseases (Vlassara & Palace, 2002; Yılmaz & Karabudak, 2016; Çetiner & Rakıcioğlu, 2020; Demirel & Yardımcı, 2022; Küçükkatırcı, Caferoğlu, & Hatipoğlu, 2022).

Since the formation of advanced glycation end products is generally due to the high amount of glucose in the medium, studies in this field have largely focused on diabetes and its complications. Although the general idea on this subject is that AGEs occur as a result of diabetes, studies show that AGEs also have diabetic effects and there are basically two mechanisms involved in the emergence of this effect. Firstly; It has been shown that AGEs can directly cause pancreatic beta-cell damage, leading to disruptions in insulin secretion. The second mechanism occurs when the accumulation of AGE on the respiratory proteins in the mitochondria induces the formation of reactive oxygen species (ROS), thus triggering a vicious inflammatory cycle (Yılmaz & Karabudak, 2016; Demirel & Yardımcı, 2022).

The role of AGEs in metabolic memory

The concept of metabolic memory; It is defined as metabolic disorders seen in various diseases, especially diabetes, and hyperglycemia that occurs even for a temporary period, mediating cardiovascular complications in the future with the permanent effect of epigenetic modifications, even if blood sugar control is achieved later on. This concept fully emphasizes the effects and complication risks of metabolic changes that occur in the early stages of pathological disorders, especially diabetes. Many researchers, who have focused on the pathogenesis of metabolic memory for years, are trying to find an answer to the question of why diabetes-related complications continue to occur in the later periods, although early glycemic control can be achieved in diabetic individuals (Küçükkatırcı, Caferoğlu, & Hatipoğlu, 2022).

One of the key factors in the emergence of diabetic complications is the production of high reactive oxygen species (ROS), which occurs in mitochondria and endothelial cells, which leads to increased oxidative stress. High amounts of glucose metabolized in mitochondria and endothelial cells appear to lead to even more free radical production. It is known that the high production of ROS in mitochondria is the first step of the chain reactions that play a role in the emergence of chronic complications of diabetes. In particular, this situation has also been described as the vicious circle of metabolic memory. In this case, free radicals released at high levels in the respiratory chain in the mitochondria; It has also been reported to trigger various intracellular pathways such as polyol and hexoamine pathway activation, protein kinase C induction and AGE formation. In addition to this situation, free radicals can also affect the expression levels of many genes that play a role in the pathogenesis of chronic diabetic complications. Protein kinase C induction; It can increase the expression levels of pro-inflammatory cytokines, various adhesion molecules and growth factors. In addition to this situation; It has been discovered that by stimulating NF- κ B, it can induce approximately 150 pro-inflammatory genes, including cyclooxygenase-2, interleukin (IL)-1, IL-6, IL-8, TNF- β and NO synthetase in vascular cells (Küçükkatırcı, Caferoğlu, & Hatipoğlu, 2022).

It is known that hyperglycemia in endothelial cells inhibits the production of NO, leading to abnormal vascular reactivation, which impairs the perfusion of tissues and organs. On the other hand, there are studies that determined that oxidative stress and high glucose levels increase inducible nitric oxide synthase (iNOS) activity and NO synthesis. This causes damage to proteins involved in mitochondrial electron transfer, decreased peroxide detoxification, and ultimately a shift of glucose metabolism to polyol and hexokinase pathways. It is observed that protein and lipid glycation, which occurs in parallel with this trend in glucose metabolism, leads to deterioration in vascular reactivation. The toxic effects on tissues and organs that develop due to glucose levels are also directly related to the increase in AGE production. Intracellular protein glycation and the resulting AGE formation affect the regulation of many genes that increase ROS production in mitochondria, as in hyperglycemia. Binding of advanced glycation end products to their receptors in various cells also leads to the induction of NF- κ B, which affects the regulation of genes associated with diabetes and its complications. It appears that increased glucose concentration in mitochondria and consequent increased AGE formation in mitochondria lead to increased ROS production and induction. In short, it is seen that the glycation of the proteins involved in respiration in mitochondria has a great importance in the concept of metabolic memory (Küçükkatırcı, Caferoğlu, & Hatipoğlu, 2022).

Retinopathy and eye complications

It has been observed that advanced glycation end products can accumulate in different parts of the eye such as the lens, cornea, Bruch's membrane, retina and optic nerve. It has been shown that this situation has a role in various complications, especially diabetic retinopathy, cataract and diabetic keratopathy in individuals with diabetes. In a study conducted on diabetic rats, compared to the control group, in addition to higher AGE accumulation in the experimental groups, there

was a higher expression of 8-hydroxyguanosine and NF- κ B in the corneal tissues, and this revealed that AGE accumulation may have a role in the pathogenesis of diabetic keratopathy. The accumulation of advanced glycation end products may also contribute to the formation of cataracts by causing accumulation in structural proteins such as tubulin, myelin and lens crystals in the eye and lens epithelial cells. In a study of individuals with type 2 diabetes, it was found that the degree of retinopathy was correlated with the levels of AGEs in samples taken from retinal blood vessels. This suggested that AGEs may have a role in the pathophysiology of retinopathy. Various studies of diabetic retinopathy; showed that it is characterized by proliferation or occlusion of intraocular blood vessels, increased permeability of capillaries, proliferative loss of endothelial cells and surrounding retinal pericytes. It has been reported that extracellular proteins modified as a result of glycation can cause retinal damage by binding to RAGE. It has been demonstrated that binding to the aforementioned RAGE receptor can induce various signaling pathways as well as increase oxidative stress. It has been observed that the interaction of advanced glycation end products with RAGE leads to an increase in ROS and induction of apoptosis in pericytes through the activation of NF- κ B and NADPH oxidase in endothelial cells. It has been determined that these events cause disruptions in the blood retinal barrier and cause mitochondrial damage to retinal neurons through ROS. In addition, various studies have shown that the interaction of AGEs with RAGE, which is formed not only in the cell but also in other parts of the eye, can potentially stimulate angiogenesis (Vlassara & Palace, 2002; Kerimoğlu & Türk, 2018; Çetiner & Rakıcıoğlu, 2020; Demirer & Yardımcı, 2022).

Effect of AGEs on renal functions

One of the most important causes of chronic kidney diseases in the world is known as diabetes; The incidence of diabetic nephropathy is approximately 30-40%. It is known that the kidneys, which are known to have a direct or indirect role in all metabolic events in the body, also have importance on AGE metabolism. It has also been shown that advanced glycation end products are absorbed through the glomerular filtrate through the proximal tubule cells and catabolized by these cells. Since the kidneys are important in AGE metabolism; In case of any damage to the kidneys, serum AGE levels are also directly increased. Elevated serum AGE levels also cause damage to the kidneys and this becomes a vicious cycle. Studies have revealed that the renal and vascular accumulation of AGEs triggers the accumulation of AGEs in most of the systems in the human body and mediates the complications that occur due to this accumulation. Nephropathy due to hyperglycemia; It is reported to be characterized by excessive ECM deposition, glomerular hypertrophy, glomerulosclerosis and thickening of the basement membrane (Vlassara & Palace, 2002; Ergenç & et al., 2017; Demirel & Yıldırım, 2018; Çetiner & Rakıcıoğlu, 2020).

It has been suggested that advanced glycation end products interact with RAGE to induce some signaling pathways and thus play a role in the development of nephropathy in diabetes. It has also been demonstrated that AGEs have a role in podocyte DNA damage and segregation, in part by stimulating the angiotensin II type 1 receptor (AT1R). Since nephropathy is characterized by the accumulation of ECM proteins in the glomerular mesangium and tubulointerstitium; It has been understood that AGEs lead to the deterioration of the balance between the synthesis and degradation of components in the ECM structure, leading to the accumulation of collagen, fibronectin and laminins, and thus to various pathologies. It has been shown that advanced glycation end products increase the formation of ECM proteins such as fibronectin and type I-IV collagen depending on dose and time. This increase was thought to be mediated directly by RAGE receptors, which involve induction of the Janus kinase (JAK)/signal converter and activator of transcription (STAT) pathways. It has been understood that these signaling pathways mediate the emergence of these effects by causing an increase in the expression of growth factors and profibrotic cytokines, including platelet-derived growth factor (TGF) and connective tissue growth factor (CTGF) (Demirer & Yardımcı, 2022).

Relationship between neuropathy and AGEs

One of the most common complications in hyperglycemia is neuropathy, and it is generally one of the most common causes of non-traumatic lower extremity amputations in individuals with diabetes. Although the pathogenesis of diabetic neuropathy is not fully understood, AGEs caused by high glucose levels are thought to have a key role in this regard. It is known that the effects of advanced glycation end products such as the polyol pathway, increase in oxidative stress and induction of the protein kinase C pathway, and that these pathways cause an increase in oxidative stress levels in general. It has been reported that oxidative stress resulting from these reasons also contributes to the accumulation of glycoxidation products such as carboxymethyllysine (CML) and pentosidine. Modification of proteins in peripheral nerves with AGEs leads to structural and functional changes. In vitro studies have shown that incubation of Schwann cells and neurons with AGEs decreases the survival rate of the cells. It has been reported that one of the main factors in peripheral nerve fiber loss seen in diabetics is AGE accumulation. The basis of this loss is thought to be the inhibition of axonal transport as a result of AGE accumulation in tubulin and neurofilaments. This inhibition of axonal transport leads to degeneration and atrophy of nerve fibers, and ultimately contributes to neuropathy. Since advanced glycation end products receptors are also found in the peripheral nervous system, previous studies have revealed that the interaction between AGEs and RAGE has various effects on the nervous system. In a study, it was shown that the AGE-RAGE interaction can induce caspase-3 activation, production of ROS, and nuclear DNA damage in dorsal root ganglion neurons. In other studies, it has been revealed that the loss of sensation in diabetic individuals and the increase in NF- κ B and NF- κ B-dependent pro-inflammatory gene expression levels in peripheral nerves are due to the interaction of AGE-RAGE. It is reported that all these pathological processes can affect every component in the peripheral nervous system cells (Demirer & Yardımcı, 2022).

Therapeutic approaches and prospects

The data obtained as a result of the current and ongoing observations obtained over the years have shown that the accumulation of AGEs in cells and tissues can cause various pathologies. In particular, studies confirm that AGEs are an important factor in the long-term complications of hyperglycemia. Therefore, it has been understood that the success of treatment options in diabetics can be achieved with early and intensive hyperglycemic control. One of the possible strategies is to try methods to reduce AGE formation, AGE receptor expression and oxidative stress, together with early and intense hyperglycemic control. One of the methods studied in this regard is the use of bioactive components such as polyphenols. These compounds; They are secondary metabolites found in the structure of plants, containing at least one carboxyl group and reported to have antiglycation activity. In various studies, it has been shown that thymoquinone, one of this group of compounds, has stronger antiglycation activity than aminoguanidine, which is an artificial antiglycation agent. It is also known that drugs that are widely used in the treatment of type 2 diabetes have an inhibitory effect on the formation of AGEs. In vitro studies with pioglitazone and metformin, which are among these drugs, have been shown to prevent AGE formation. Angiotensin converting enzyme (ACE) inhibitors and angiotensin II subtype 1 receptor (AT-1) blockers have been found to be able to reduce AGE formation while providing blood pressure control. Telmisartan, on the other hand, is a compound that has been shown to be able to gradually inhibit the formation of superoxide over the years, as well as reduce the levels of AGE receptor messenger RNA (mRNA). In addition, these drugs also have antioxidant effects. There is evidence that especially AT-1 blockers are directly effective against oxidative stress resulting from hyperglycemia and have a role in metabolic memory. Considering all these data together, it is clear that mixed therapies should be preferred, which includes methods that prevent AGE formation while treating hyperglycemia in treatment approaches. Preferring such combined methods in treatment; It is quite clear that it will make significant contributions to the prevention of many diabetes-related microvascular and peripheral - cerebrovascular diseases such as nephropathy,

retinopathy and neuropathy, and macrovascular complications related to diabetes such as ischemic heart disease and hypertension (Küçükkatırcı, Caferoğlu, & Hatipoğlu, 2022; Yalçın & Rakıcıoğlu, 2022).

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Non-Arteritic Anterior Ischemic Optic Neuropathy Developing after COVID-19 mRNA Vaccine

Ramazan Birgöl¹

Introduction

The COVID-19 virus, which emerged in China in 2019, is affecting the entire world, and mankind has mobilized all its means for such a mortal virus. Vaccination, which is the most effective way, is among them. One of the most widely used vaccine types in the world is the BioNTech mRNA COVID-19 vaccine. However, the vaccine, which is widely used all over the world, is likely to have side effects.

Non-arteritic anterior ischemic optic neuropathy (NAAION) occurs with a sudden and dramatic decrease in vision. Its clinical manifestation takes the form of a swollen and edematous optic disc. There are various diseases, such as diabetes mellitus (DM), hypertension, anemia and hyperlipidemia, which are among the most common causes of NAAION.^{1,2}

Radiological, biochemical and microbiological examinations were carried out for the patient in this case. No significant evidence was detected for the development of NAAION for the aforementioned reasons. The patient specifically stated that the clinical manifestation of the patient developed in a very short time after receiving vaccine (BioNTech, Pfizer). Despite the frequent administration of the COVID-19 mRNA vaccine (BioNTech, Pfizer), no cases with optic neuropathy involvement were reported in the literature after receiving the COVID-19 mRNA vaccine (BioNTech, Pfizer). This case is the first to develop NAAION following the COVID-19 mRNA vaccine (BioNTech, Pfizer).

The Case

A 53-year-old female patient was admitted to our clinic with sudden vision loss in the left eye. The patient suddenly noticed a narrowing in the lower-left visual field three days after the second dose of the BioNTech mRNA COVID-19 vaccine. She also experienced fatigue and muscle and joint pain, which are common side effects after the vaccination. It was recorded that the patient had diabetes as a comorbidity and was taking metformin. The patient stated that her diabetes was under control. No smoking or alcohol intake was present.

The patient's vision was 1.0 with correction on the right and at hand movement level on the left. The anterior segment was normal bilaterally. Intraocular pressure was 14 mmHg on the right and 16 mmHg on the left. There was a relative afferent pupillary defect on the left. There was no pain with eye movements. After the patient's pupil was dilated, the right fundus was normal, and the optic disc was swollen and edematous in the left fundus. Optical coherence tomography (OCT), fundus fluorescein angiography (FFA), and cranial and orbital magnetic resonance (MR) were performed for the patient, who was hospitalized and received pulse steroid therapy at 1 g a day for three days. The steroid treatment was continued orally for 14 days and was terminated by

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lowering the dose. The sedimentation, CRP and whole blood levels were checked before the steroids, and they were found to be within normal values. Biochemical and microbiological parameters, such as fibrinogen, homocysteine, the Epstein Barr Virus, c-ANCA, p-ANCA, TORCH panel and adenosine deaminase levels were checked to rule out the causes of NAAION. Neurology and internal medicine department consultations were requested from the patient. No systemic condition was detected in the patient's blood parameters, and MRIs were taken to explain this during all this time. The OCT, fundus images, FFA, visual field, MRI examination results and blood parameters of the patient are shown below (Figure 1a, 1b, 1c , 1d, 1e, 1f).

Figure 1a. Fundus images of the patient

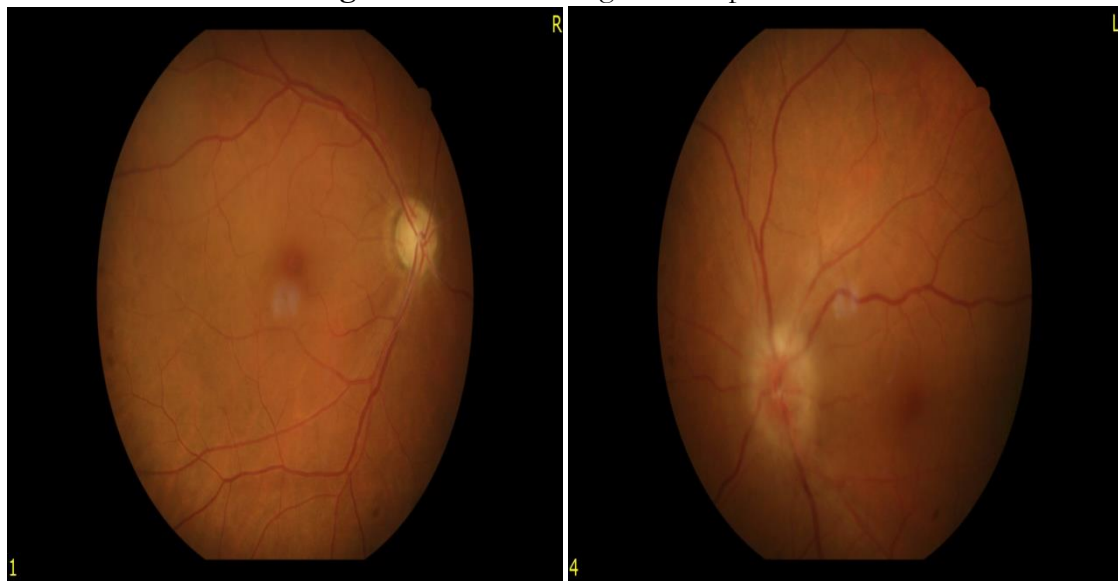
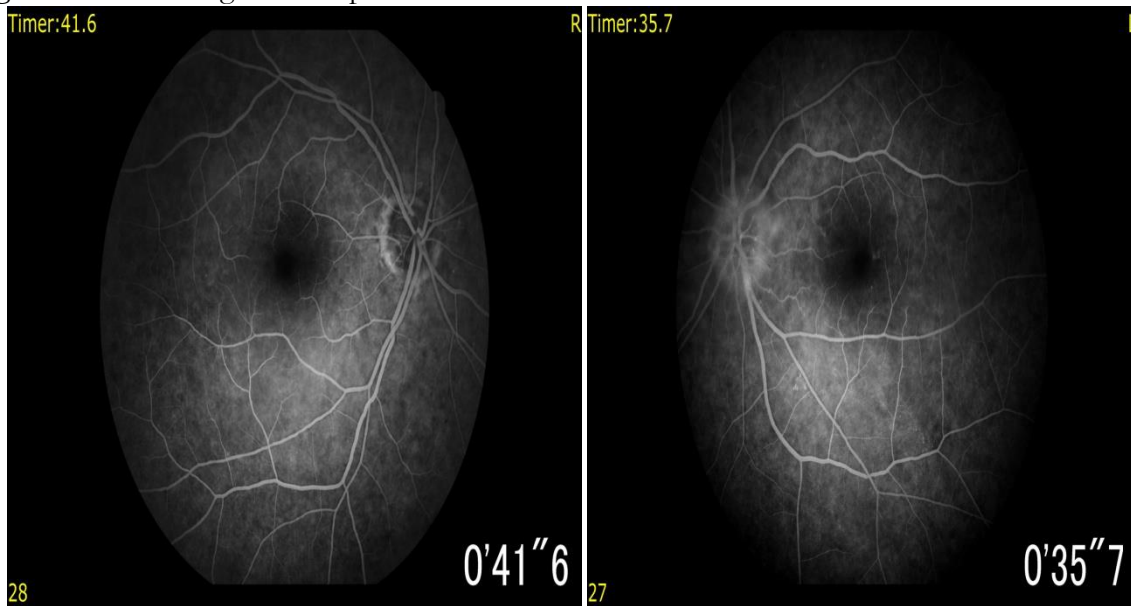


Figure 1b. FFA images of the patient



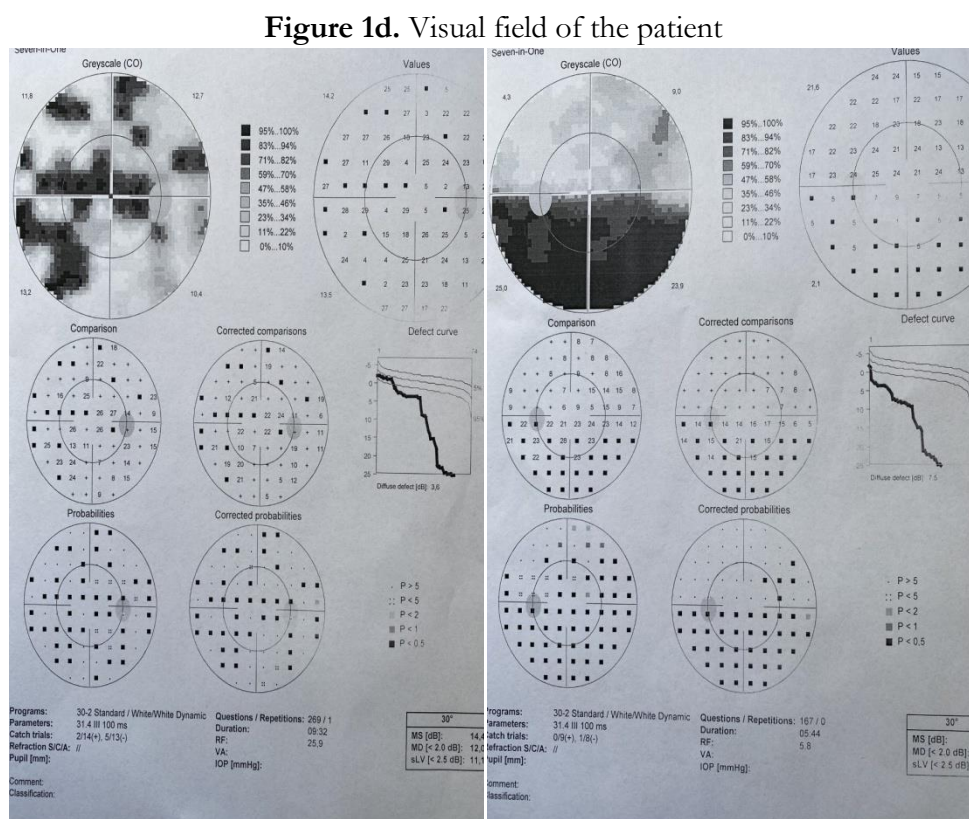
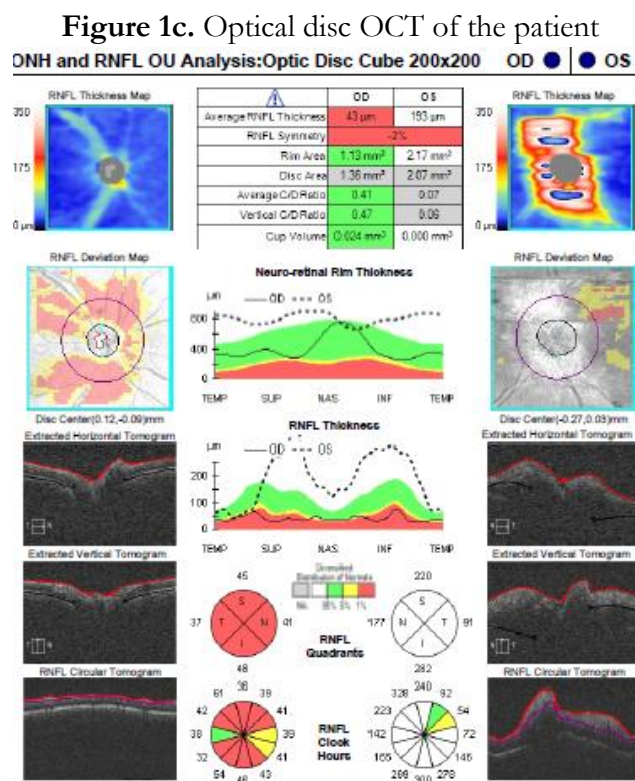


Figure 1e. Orbital MRI of the patient

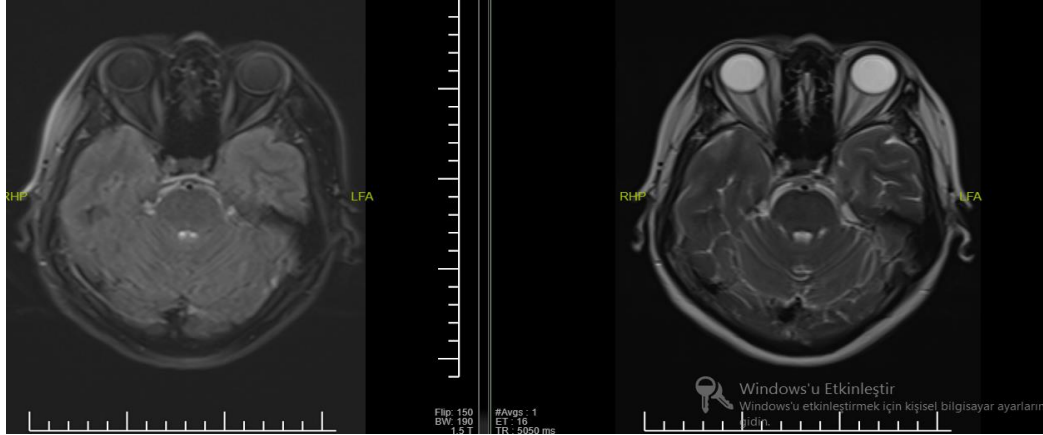


Figure 1f. Blood parameters of the patient

Tetkik Adı	Sonuç	Duru	Birim	Referans Aralığı
WBC	7,79		10 ³ /μL	4 - 10,5
NEU	4,63		10 ³ /μL	2 - 7
NEU%	59,4		%	40 - 70
LY%	34,1		%	20 - 50
LYM	2,65		10 ³ /μL	0,8 - 4
EOS	0,14		10 ³ /μL	0,02 - 0,5
EO%	1,8		%	0,5 - 5
BASO	0,03		10 ³ /μL	0 - 1
BASO%	0,3		%	0 - 1
MONO	0,34		10 ³ /μL	0,12 - 1,2
MON%	4,4		%	3 - 12
RBC	4,72		10 ⁶ /μL	3,5 - 5
HGB	12,6		g/dL	12 - 16
HCT	38,0		%	37 - 47
MCV	80,5		fL	80 - 100
MCH	26,7	(D)	pg	27 - 34
MCHC	33,2		g/dL	32 - 36
RDW-CV	13,1		%	11 - 16
PLT	156		10 ³ /uL	150 - 450
MPV	10,9		fL	6,5 - 12
PCT	0,169		%	0,108 - 0,282
PDW	16,5		%	10 - 65
Sedimentasyon	13		mm	0 - 20
GGT	133	(Y)	IU/L	9 - 36
Total Protein	71,3		g/L	63 - 86
Albumin	46		g/L	35 - 55
Bilirubin (direkt)	0,25		mg/dL	0 - 0,5
Bilirubin (indirekt)	0,06	(D)	mg/dl	0,1 - 1
Bilirubin (total)	0,31		mg/dL	0 - 1
Kalsiyum (Ca)	9,2		mg/dL	8,4 - 10,2
Potasyum	3,8		mmol/L	3,5 - 5,1
Sodyum	146		mmol/L	136 - 146
Klor(Cl)	106		mmol/L	96 - 110
CRP	1,7		mg/L	0,1 - 5
Ürik asit	5,0		mg/dL	2,3 - 8,2
Glukoz (açlık)	148	(Y)	mg/dL	70 - 105
BUN	12,0		mg/dl	10 - 20
Kreatinin	0,71		mg/dl	0,6 - 1,1
t GFH (CKD-EPI formülü ile)	>90		-	-
CK	51		U/L	29 - 200
CK-MB	9,3	(D)	U/L	10 - 25
AST	43	(Y)	U/L	5 - 34
ALT	56	(Y)	U/L	0 - 55
ALP	93		U/L	40 - 150
HDL kolesterol	48,8		mg/dL	35 - 60
Trigliserid	153		mg/dL	50 - 200
LDL kolesterol (Hesaplamalı)	112,6		mg/dL	60 - 130
Kolesterol	192		mg/dL	0 - 200

Serbest T3	3.08		ng/L	2.3 - 4.2
Serbest T4	1.31		ng/dL	0.85 - 1.6
TSH	0.269	(D)	mIU/L	0.55 - 4.78
Vitamin B12	356		ng/L	214 - 914
Anti HBs	0.37		mIU/mL	0 - 9.9
Anti HCV	0.40		mIU/mL	0 - 0.99
Anti HIV	0.05		mIU/mL	0 - 0.99
Anti CMV IgM	0.14		mIU/mL	0 - 0.99
Anti CMV IgG	141.6	(Y)	AU/mL	0 - 5.9
VDRL-RPR	Negatif (-)	-	-	-
EBV EBNA IgG	15.78		-	-
EBV VCA IgG	41.02		-	-
EBV VCA IgM	0.10		-	-
Anti Rubella IgM	0.27			0 - 1.2
Anti Rubella IgG	37.4	(Y)	IU/mL	0 - 4.9
Anti Toxoplasma IgM	0.21			0 - 0.49
Anti Toxoplasma IgG	6.3	(Y)	IU/mL	0 - 1.6
HLA B27	NEGATİF	-	-	-
p-ANCA (antimiyeloperoksidaz) (ELISA)	1.15 NEGATİF		U/mL	-
Adenozin deaminaz aktivitesi (ADA)	16.86		U/L	5.0 - 20.0
c-ANCA (antimiyeloperoksidaz) (ELISA)	0.33 NEGATİF		U/mL	-
Homosistein	9.31		μmol/L	4.65 - 12.22
Anti nükleer antikor (ANA)	NEGATİF	-	-	NEGATİF -

Discussion

There is no doubt that the vaccine plays one of the biggest roles in the effort to stop the COVID-19 pandemic, which has affected the entire world. Most vaccines are safe, but they usually have a wide spectrum of side effects, and it is equally difficult to prove that the side effects are related to the vaccine.³ It is reported in the literature that vaccines generally cause side effects through the adjuvants they include, which can happen in two ways. They cause these side effects either by initiating an inflammatory event or by triggering an autoimmune mechanism.⁴ We believe that an autoimmune mechanism played a role in this case.

It was reported that central serous chorioretinopathy developed in only one patient as an intraocular complication after receiving the COVID-19 mRNA vaccine (BioNTech, Pfizer).⁵ Optic neuropathy development has not been reported in the literature. It was detected that optic neuropathy develops after vaccinations such as influenza, anthrax and rubella. This situation, which may be unilateral as reported in some publications, can sometimes be bilateral. Treatments were generally initiated with pulse steroids and continued with oral treatment for 14 days. Although visual acuity increased in some cases, it generally remained the same or deteriorated.^{6,7} We applied the same treatment modality in this case, and the patient's vision remained at the same level and did not improve.

This case is very important in terms of the immediate development after the vaccination. Furthermore, the fact that NAAION is more common in men, and the fact that the age of consent was 60, further distinguishes this case from classical causes of NAAION.⁸ DM, which could cause possible NAAION in the patient, was ruled out as a result of the long-term regulation of her blood sugar and internal medicine consultations. Neurological causes were also ruled out as a result of the absence of significant pathologies in cranial and orbital MRIs and neurology consultations. All of the radiological, biochemical and microbiological results indicated that the most likely cause of NAAION was vaccination.

In conclusion, COVID-19 vaccinations, which are administered all over the world, naturally increase the incidence of side effects. This case will contribute to the literature because it is the first case that developed NAAION following the COVID-19 mRNA vaccine (BioNTech, Pfizer).

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Current Approach to Molecular Mechanism pf Antibiotic Resistance in Enterobacteriaceae

Seyda İĞNAK TARLIĞ

Enterobacteriaceae is a family of gram-negative rods commonly found in nature and intestinal flora of most animals, including humans. Enterobacteriaceae is responsible for clinically significant health-care and community acquired infections in human. β -lactam group drugs are used extensively in the treatment of Enterobacteriaceae-related infections. It is known that the overuse of these antibiotics promotes the problem of antibiotic resistance. Today, bacteria that are members of the Enterobacteriaceae can develop resistance to broad-spectrum beta-lactam antibiotics and carbapenems with inactivating enzymes called beta-lactamases. This does not only cause problems in the treatment of Enterobacteriaceae-related infections, but also it increases mortality rates, duration of hospital stay, and health care costs all over the world (Murray, Rosenthal & Pfaller, 2021, De Angelis et al., 2020).

Resistance genes encoded in mobile genetic elements such as transposon, integron, and plasmid cause the spread of antibiotic resistance among different bacterial species. The World Health Organization (WHO) states that multi-drug-resistant Gram-negative bacteria (MDR-GNB), including carbapenem-resistant Enterobacteriaceae (CRE), have priority in new antibiotic research and development. The reason for this high priority is the necessity of infection control measures to reduce the spread of infection in the community and health care services, and the high mortality observed especially in serious infections in immunocompromised individuals (WHO, 2017). Centers for Disease Control and Prevention (CDC) emphasizes that carbapenem-resistant Enterobacteriaceae is a global threat due to the use of toxic and less effective methods in treatment due to their resistance to almost all antibiotics. (CDC, 2019).

Enterobacteriaceae

Members of the Enterobacteriaceae family are gram-negative rods of 0.3-1.0 x 1.0-6.0 μm in size, which do not form spores. Enterobacteriaceae, which have simple nutritional requirements, ferment glucose, reduce nitrate, are catalase positive and oxidase negative. It is commonly found in the intestinal flora of many animals, including humans, and in nature such as soil, water and vegetation, especially in humid environments.

Some Enterobacteriaceae members (*Salmonella typhi*, *Shigella dysenteriae*, *Yersinia pestis*) are obligate pathogens and their detection in clinical samples is always associated with infection, while some (*Escherichia coli*, *Klebsiella pneumoniae* etc.) are opportunistic pathogens which cause serious infections, especially in immunocompromised individuals (Murray, Rosenthal & Pfaller, 2021).

Enterobacteriaceae may cause community-acquired, hospital-acquired, and health care-associated infections. Enterobacteriaceae strains which are isolated from rectal swab, urine and respiratory tract samples may be associated with colonization rather than infection. The inability to distinguish between colonization and infection easily causes trouble in treating patients, especially in areas where multidrug resistance is common (Paterson & Mathers, 2020).

Mechanisms of Antibiotic Resistance

Antibiotic resistance in microorganisms can be classified into two as natural and acquired resistance.

1) Natural (Intrinsic) Resistance: It depends on the natural characteristics of microorganisms encoded on the chromosome. This type of resistance is independent of the selective pressure of antibiotics and horizontal gene transfer.

Natural resistance can develop with different characteristics encoded on the chromosome such as lack of a target of the drug, not taking the drug into the cell by the bacteria, removal of the drug from the cell by the efflux pump systems, and the presence of enzymes that inactivate the drug. Knowing the natural resistance profiles of microorganisms will prevent unnecessary antibiotic use and thus the development of acquired resistance. For example, *Klebsiella pneumoniae* is naturally resistant to ampicillin. This resistance is thought to be caused by the β -lactamase enzyme, which is a product of the SHV-1 gene encoded on the chromosome. As a result of the intensive use of ampicillin and antibiotics in the same spectrum as ampicillin in hospital settings, *K. pneumoniae* colonization is found in the intestinal flora of the patients. Prolongation of hospital stay and antibiotic use increase the importance of *K. pneumoniae* in nosocomial infections (Patterson et al., 2000, Paterson & Mathers, 2020, Chaves et al., 2001).

2) Acquired resistance: Acquired resistance can occur as a result of mutation in an existing gene or by the acquisition of new resistance genes through horizontal gene transfer mechanisms such as conjugation, transformation and transduction. Contrary to natural antimicrobial resistance, it first appears in some strains and subpopulations of the bacterial species. The intracellular concentration of the antibiotic can not reach the desired therapeutic concentration due to decreased uptake (down-regulation of porin protein expression, etc.) or increased multi-drug resistance pumps, changes in the target of the drug through mechanisms such as mutation or post translational modification, and enzymatic inactivation of the antibiotic (Paterson & Mathers, 2020, Opal & Pop-Vicas, 2020).

Molecular Mechanisms of Resistance in Enterobacteriaceae

Enzymatic Inhibition of Antimicrobial Activity

Bacteria may gain antimicrobial resistance by breaking down the antibiotic with various enzymes or by impairing their antibacterial activity. There are different types of enzymes with hydrolyzing (β -lactamase), inactivating (aminoglycoside-inactivating enzymes, chloramphenicol acetyltransferase, erythromycin esterase etc.) and modifying (aminoglycoside modifying enzyme) types of activities (Zhu et al., 2022, Opal & Pop-Vicas, 2020).

Enzymatic inactivation of β -lactam antibiotics

Since the discovery of benzylpenicillin in the 1920s, many penicillin derivatives and β -lactam group antibiotics such as cephalosporins, monobactams and carbapenems have been discovered. These bactericidal drugs inhibit bacterial cell wall biosynthesis by covalently binding to transpeptidases (Penicillin Binding Proteins-PBPs) (Bush & Bradford, 2016).

Resistance to β -lactam antibiotics is mainly caused by β -lactamase enzyme that cleaves the amide bond in the β -lactam ring of the antibiotic. This resistance mechanism probably developed by bacteria against natural antimicrobials and has become common due to increased use of antimicrobial drugs in modern medicine.

The genes encoding the β -lactamase enzyme can be encoded on the chromosome or on mobile genetic elements such as plasmid and transposon. It has also been shown that β -lactamase genes can be encoded on the integron. According to the Ambler classification (Ambler, 1980), β -

lactamases are classified based on their amino acid sequence into 4 molecular groups (A, B, C and D). Class B beta-lactamases called metallo- β -lactamases break the amide bond in the beta-lactam ring using zinc (Zn^{2+}), whereas class A, C and D beta-lactamases hydrolyze the beta-lactam ring by using the serine residues in their active sites (De Angelis et al., 2020, Opal & Pop-Vicas, 2020).

Most of the extended-spectrum beta-lactamases are in Ambler group A and are inhibited by clavulonic acid and tazobactam (Zhu et al., 2022). Ampicillin resistance in Gram-negative bacteria increased in 1960s with TEM-1 (Temoneria β -lactamase) beta-lactamase encoded on the plasmid. SHV type β -lactamase associated with TEM-1 in terms of molecular structure has become common especially in *E. coli* and *K. pneumoniae* isolates. TEM-1 is the most common β -lactamase in Gram-negative bacteria, hydrolyzes penicillin and narrow-spectrum cephalosporins in the Enterobacteriaceae. The configuration change of a few amino acids in the active site of the TEM-related ESBL enzyme increases the effect of the enzyme against 3rd generation cephalosporins (cefotaxime, cefpodoxime, ceftazidime, ceftriaxone) and monobactam (aztreonam). The first TEM-related ESBL (TEM-3) was reported in 1988. Upto date there are more than 200 TEM-related ESBL resistances identified. These enzymes are mostly observed in members of Enterobacteriaceae such as *E. coli* and *K. pneumoniae*. SHV-1 type β -lactamase is mainly found in *K. pneumoniae* strains. Point mutations in the active site of SHV-1 β -lactamase, which has 68% similarity to TEM-1 at amino acid level, has been caused the development of SHV type ESBL (Opal & Pop-Vicas, 2020). Some specific OXA β -lactamases (OXA-10, OXA-13, OXA-19) are included in the ESBL group because they hydrolyze extended spectrum cephalosporins (Zhu et al., 2022).

Cefotaxime-M (CTX-M) enzyme has spread rapidly around the world and has become among the most common ESBL in the world. In general, CTX-M hydrolyzes cefotaxime and ceftriaxone better than ceftazidime and is inhibited more by tazobactam than clavulanic acid (Opal & Pop-Vicas, 2020). The ST131 clone carrying the CTX-M-15 gene encoded on the plasmid plays an important role in multidrug resistant *E. coli*-associated infections in Europe and America (Johnson et al., 2010). CTX-M-14 is commonly detected in South East Asia whereas CTX-M-27 in Japan and Europe (Zhu et al., 2022).

AmpC β -lactamases confer resistance to penicillin, narrow-spectrum cephalosporins, oxymino-beta-lactams and cephamycins, but they are not sensitive to beta lactamase inhibitors such as clavulanic acid and tazobactam. While they do not show any effect against cefepime and aztreonam, changes in the active sites with point mutations may increase their effects on cefepime. Although AmpC β -lactamases are listed in the Ambler C group, there are also several members in the A group. They can be encoded on the chromosome or plasmid. Low level AmpC production is observed in *E. coli* strains while there is no AmpC coding gene on the chromosome of *Klebsiella* and *Salmonella* strains. But these bacteria may acquire AmpC genes which are encoded on the plasmids (Opal & Pop-Vicas, 2020, Zhu et al., 2022). Although their production is normally suppressed in Gram-negative rods, inducible Amp-C production may be observed after exposure to β -lactam antibiotics. Amp-C beta lactamase production decreases in a short time after the antibiotic exposure is terminated. However, irreversible overproduction of Amp-C may be seen due to point mutations (Opal & Pop-Vicas, 2020). AmpC enzymes encoded on the plasmid are named according to the drug that provide resistance (CMY, FOX, MOX, LAT), enzyme type (AAC, ACT) or the region they were discovered (MIR-1 or DHA) (Zhu et al., 2022).

Enzymatic inactivation of carbapenem antibiotics

Carbapenemases are β -lactamase enzymes that hydrolyze not only carbapenems but also broad-spectrum penicillins, oxymino-cephalosporins, and cephamycins. Enterobacteriaceae develop resistance by five types of carbapenemases such as *Klebsiella pneumoniae* carbapenemase (KPC), New-Delhi metallo-beta-lactamase (NDM), imipenem's metallo-beta-lactamase (IMP),

Verona integron encoded metallo- β -lactamase (VIM) and oxacillin carbapenemase (OXA) that are encoded on mobile genetic elements (Opal & Pop-Vicas, 2020).

Although KPC, an Ambler class A serine carbapenemase, was first identified in *K. pneumoniae* strains, it is now commonly detected in *E. coli*, *Citrobacter*, *Enterobacter*, *Salmonella* and *Serratia* strains worldwide (Logan & Weinstein, 2017).

Ambler Class B metallo- β -lactamases (MBLs), which are encoded on the chromosome and break the beta-lactam ring using Zn^{2+} , were mostly detected in environmental bacteria with low pathogenic potential. Most clinically important transmissible carbapenemases are in NDM, IMP and VIM. NDM-1 was first detected in *K. pneumoniae* strains in India in 2008, it is now widely observed in many countries because of rapid gene transfer between species (Opal & Pop-Vicas, 2020, Logan & Weinstein, 2017).

OXA-48-like carbapenemases belong to Ambler Class D. Among these variants OXA-48, which is increasingly identified in Enterobacteriaceae, provides weak but significant carbapenemase activity (Poirel, Potron & Nordman, 2012).

Enzymatic inactivation of aminoglycoside antibiotics

Resistance to aminoglycoside antibiotics usually develop through aminoglycoside-modifying enzymes encoded on chromosomes or mobile genetic elements. These enzymes modify aminoglycoside antibiotics by phosphorylation of the OH group, adenylation of the OH group (addition of adenosine monophosphate-AMP), or acetylation of the NH₂ group (Clark et al., 2019). According to their modifying properties and activities, they are divided into three groups: acetyltransferases (AAC), nucleotidyl transferases (ANT), and phosphotransferases (APH) (Ramirez & Tolmasky, 2010). There are many enzymes that target specific amino and hydroxyl groups for a activity. Of these, aminoglycoside acetyltransferase (AAC) acting in the 3' region is named AAC(3'), while there is more than one enzyme catalyzing the same reaction, it is named with Roman numerals (For example, AAC(3')-IV). The distribution of aminoglycoside resistance genes in the world is related to the selective pressure of the commonly used antibiotic. APH(3'), which causes decreased susceptibility to kanamycin and streptomycin, spreads in Gram-positive and negative bacteria worldwide.

Aminoglycoside nucleotidyltransferase (ANT)(2'') has been associated with multiple nosocomial outbreaks in the United States in 1990. It was determined that *K. pneumonia* strains carrying ANT(2') gene encoded on plasmid are responsible for a significant portion of the outbreaks associated with aminoglycoside resistant Enterobacteriaceae. During epidemics, the plasmid ANT(2') gene was observed to be spread in other strains of the species and Enterobacteriaceae member bacteria

AAC(6') is common in enteric bacteria in East Asia. AAC(3') group enzymes were seen in many epidemics in South America, Western Europe and the United States. (Opal & Pop-Vicas, 2020).

The first bifunctional enzyme (AAC(6')-Ib-cr) that inactivates aminoglycosides and fluoroquinolone (ciprofloxacin) was discovered in 2006 (Robicsek et al., 2006).

Enzymatic inactivation of tetracycline antibiotics

Although tetracycline resistance occurs through mechanisms such as changes in the ribosomal target or efflux pump systems, it is expected that the tetracycline-degrading enzymes called TetX which are seen in environmental bacteria may also be transmissible to strains with clinical importance (Forsberg, 2015). In 2013, the plasmid-encoded tetX gene was shown in Enterobacteriaceae strain, which was the causative agent of urinary tract infection for the first time (Leski & et al., 2013).

Enzymatic inactivation of Macrolide-Lincosamide-Streptogramin antibiotics

Resistance to erythromycin and other antibiotics develops as a change in the target of the drug, removal of the drug from the cell by efflux pump systems, and inactivation of the drug by enzymes. Erythromycin esterase, one of the inactivating enzymes, hydrolyzes the lactone ring of erythromycin. Erythromycin esterase encoded on the plasmid causes high levels of erythromycin resistance. (Ounishi & Courvalin, 1985, Arthur, Autissier & Courvalin, 1986, Opal & Pop-Vicas, 2020). In addition, erythromycin esterase gene that causes high-level macrolide resistance and NDM-1 type carbapenemase were detected in a same *K. pneumoniae* strain (Yong et al., 2009).

Decreased permeability of bacterial membrane

Diffusion channels called porins in the cell membrane are used for hydrophilic antibiotic molecules to pass through the outer cell membrane. Numerous porins (10^5 porin molecules per one *E. coli*) are found on bacterial membranes. Bacteria change the porin number according to the osmolarity of the medium. In the hyperosmolar environment, *E. coli* decreases the expression of larger porins (OmpF) while maintaining the production of small porins (OmpC). The passage of antibiotics across the cell membrane is not only related to the number and structure of porin proteins, but also to the physiochemical structure of the antibiotic. It is difficult for negatively charged, hydrophobic and large antibiotic molecules to pass through porins. While the cell membrane is permeable to small, hydrophilic and zwitterionic charged antibiotic molecules such as imipenem, it is not permeable to large and highly charged molecules such as carbenicillin (Opal & Pop-Vicas, 2020). The number of OmpF porin proteins is decreased in *E. coli* strains via chromosomal Mar mutation. It is known that decreased OmpF porin expression, as in the marA mutation, causes resistance to beta-lactams, tetracycline, chloramphenicol and hydrophilic fluoroquinolones (Cohen, McMurry & Levy, 1988). Mutations in genes encoding porin protein are known to cause increased resistance to β -lactam antibiotics. Loss of porin proteins (OmpK35, OmpK36) in *K. pneumoniae* strains has been shown to increase carbapenem resistance in strains carrying ESBLs (CTX-M or SHV) or carrying AmpC (ACT-1, CMY-2, CMY-4 or DHA-1) enzymes which are encoded on plasmids (Crowley, Benedi', & Dome' nech-Sa'nchez, 2002, Lee et al. 2007).

In addition, it was determined that resistance developed against aminoglycoside and carbapenem group antibiotics due to the decrease in outer membrane proteins during treatment. Plasmid-encoded chloramphenicol resistance has been demonstrated in *E. coli* strains due to decreased membrane permeability (Gaffney, Cundliffe & Foster, 1981).

Aminoglycoside antibiotics are taken into the cell by anionic transporters. Positively charged antibiotics can pass through the cytoplasmic membrane due to negatively charged intracellular environment. Before this transition is achieved, a threshold value of negative charge must be present in the cell. In some cases where the proton motive force or energy production is mutated, resistance to aminoglycosides may develop. Aminoglycoside resistant mutants due to electron transport system defects have been observed in *E. coli*, *Salmonella spp.* (Opal & Pop-Vicas, 2020).

Antibiotic efflux pumps

The extrusion of the toxic material such as antibiotics out of the cell by the energy dependent pump systems causes antibiotic resistance by reducing the intracellular concentration of the drug. There are five major subfamilies of efflux pumps; 1) Resistance Nodulation Division (RND) family, 2) Major Facilitator Superfamily (MFS), 3) ATP binding cassette (ABC) superfamily, 4) Small Multidrug Resistance (SMR) family, 5) Multidrug and Toxic Compound Extrusion (MATE) (Thakur V, Uniyal A & Tiwari V, 2021).

The extrusion of the antibiotics out of the cell by the energy dependent pump systems causes resistance by reducing the intracellular concentration of the drug. It is known that AcrAB-TolC

and OqxAB-TolC efflux pumps, which are members of the RND family, cause resistance to fluoroquinolones, β -lactams, novobiocin, tetracycline, erythromycin, linezolid, quinolone, chloramphenicol, nitrofurantoin in members of Enterobacteriaceae (Thakur V, Uniyal A & Tiwari V, 2021).

It is known that EmrAB-TolC, a member of the MFS type efflux family, provides resistance to novobiocin and fluoroquinolones, while MacAB-TolC, a member of the ABC efflux family, causes macrolide resistance in Enterobacteriaceae (Nishino, Latifi & Groisman, 2006, Kobayashi, Nishino & Yamaguchi, 2001).

While QepA causes low-level fluoroquinolone resistance, it is thought that it may be responsible for the development of high resistance when combined with resistance mechanisms such as DNA gyrase mutations. (Yamane et al., 2007).

There are over 40 identified tetracycline resistance determinants that have been associated with drug efflux. Tet(A) and Tet(B) are the common tetracycline efflux pumps in Gram negative bacteria (Chopra & Roberts, 2001, Grossman, 2016).

Changes in antibiotic targets

The effect of the drug can be prevented by changes made in the target site to which the antibiotic binds. In Enterobacteriaceae, resistance to aminoglycoside antibiotics can develop by modifying the specific sites in 16S rRNA by methyl-transferases. It is known that some genes (armA, rmtA, rmtB, rmtC, rmtD, rmtE and npmA) encoding these methylase enzymes are carried on the plasmids. (Wachino & Arakawa, 2012).

Chloramphenicol inhibits the peptidyl transferase by binding to 23S rRNA which is in the large subunit of the ribosome. Resistance to chloramphenicol in Gram-negative bacteria is generally provided by the enzyme chloramphenicol acetyltransferases (CAT). This enzyme is encoded on the plasmid or chromosome and inactivates the antibiotic by O-acetylation of the 3-hydroxyl group (Clark, Pazdernik & McGehee, 2019).

DNA gyrase (Topoisomerase II) is involved in the super-folding process of chromosomal DNA during cell division. Another enzyme, topoisomerase IV, is involved in the segregation of the bacterial genome in two new cells after cell division. These enzymes consist of 2 subunits, A and B. The A unit is encoded by the *gyrA* gene, while the B region is encoded by *gyrB*. In enteric bacteria, the *gyrA* mutation is the most common cause of fluoroquinolone resistance. Plasmid-encoded quinolone resistance has been demonstrated in various members of Enterobacteriaceae. This type of resistance develops by the binding of *qnr* gene (*QnrA*, *QnrB*, *QnrS*) products to DNA gyrase. Plasmid-encoded *Qnr* genes associated with fluoroquinolone resistance low level fluoroquinolone resistance. However, because the same plasmids may encode the different genes required for the development of resistance to other antibiotics, the problem of multi-drug resistance arises in the clinic (Opal & Pop-Vicas, 2020, Nordman & Poirel, 2005).

Polymyxins, which are cationic peptide antibiotics, change the outer membrane permeability electrostatically by binding to the anionic lipopolysaccharide in the outer membrane in Gram-negative bacteria. Polymyxin B (colistin) is one of the limited number of alternatives in the treatment of multidrug resistant bacteria. Membrane modification by adding cationic groups (4-amino-4-deoxy-L-arabinose (L-Ara4N) and phosphoethanolamine (pEtN) to the cell membrane reduces the affinity of the drug to its target. Many operons and gene regions responsible for cationic modification of the cell membrane are identified. The most important ones are the dual PhoPQ and PmrAB system. Chromosomal *eptA* or *mcr* genes are carried on plasmids cause polymyxin resistance by adding pEtN to lipid A with phosphoethanolamine transferase enzyme. It is known that *mcr1-3* genes carried in plasmid spread rapidly in enteric bacteria (Opal & Pop-Vicas, 2020, Hugry et al., 2020).

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New Treatments For Primary Teeth With Excessive Substance Loss

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Introduction:

In poor nations, dental caries is a health issue that affects 60–90% of those with low income levels (Petersen, P. E., & Lennon, M. A. 2004). The breakdown of the enamel-dentin tissues of teeth by the acids created by the fermentation of the carbohydrates ingested by bacteria is known as tooth decay (Margolis and Moreno 1994). Because the structure of the enamel in deciduous teeth differs from that of permanent teeth, caries in deciduous teeth advances more quickly than in permanent teeth (Goldberg, M 2017).

Treatment of caries provides very important benefits such as cleaning the decayed tissues on the tooth surfaces, eliminating the areas where new caries may occur, stopping demineralization, restoring the tooth structures in terms of function, stopping the spread of infection, and preventing tooth loss (Downer et al. 1999; Lenters, van Amerongen, and Mandari 2006). In the article published by Hickel et al. in 2006, it was stated that the probability of failure of restorative treatments in primary teeth is higher than that in permanent teeth. In addition, the falling times of the teeth for which restorations will be made should be considered. In the guide published by the American Academy of Pediatric Dentistry (AAPD) in 2019, two general recommendations were made regarding when dental caries should be restored.

First, the risk of progression and prognosis of the caries should be evaluated specifically for that patient, and the preventive treatments should be supported by restorative treatments. The second recommendation is that the caries lesions should be restored in cases where the enamel cavitation is clinically detected, the caries image reflects from under the enamel, and caries are diagnosed in the radiographic examination ("AAPD | Pediatric Restorative Dentistry" n.d. 2022).

Primary tooth decay that is left untreated spreads quickly and results in material loss. Complex procedures can be used to treat teeth that have caries on many surfaces. Primary teeth may have more failures in caries treatment than permanent teeth. These issues can be attributed to a variety of factors, including improper material use, difficulties with dental hygiene in youngsters, tooth morphological variations, and variations in the chemical composition of teeth.

There are studies and guides showing that covering the entire deciduous molar is more successful in teeth with excessive material loss, such as those with multiple caries, pulp amputation, or root canal treatment ("AAPD | Pediatric Restorative Dentistry," n.d., 2022; Sonmez and Duruturk 2010). In order for the entire tooth to be covered with prefabricated crowns to give a successful result, it must have a good edge fit and be done in an appropriate way that will not allow plaque accumulation (Sharaf and Farsi 2004). It was reported in a study conducted by Belduz Kara and Yilmaz in 2014 that SCC types that can be made can cause gingivitis, even though they provide suitable conditions for gingival health ("Beldüz Kara, N., & Yilmaz, Y. 2014).

The American Academy of Pediatric Dentists (AAPD) suggests stainless steel crowns (PSCC) for the repair of teeth with numerous surface cavities. ("AAPD | Pediatric Restorative Dentistry," n.d.2022) Additionally, in teeth with significant substance loss, glass ionomer cements (CIS), amalgam, compomer, strip crowns, polycarbonate crowns, and zirconium crowns are also preferred. When compared to one another, these materials differ in their advantages and weaknesses. The material a dentist chooses should be unique to each instance and should take into

account cost, durability, conservatism, allergic reactions, and aesthetics. Currently, teeth with appealing appearances and socio-economic. It is regarded as a sign of development (Srinath 2017).

Until permanent teeth erupt, milk teeth perform the functions of phonation, aesthetics, and space holding. Therefore, it is crucial that the deciduous tooth remain in the mouth until it is time to fall out in order for the jaw and permanent dentition to develop properly. (Pinkham JR et al. 2005) Although the same methods are used to restore primary teeth as permanent teeth, morphological and histological distinctions can occasionally lead to divergent preferences. The need for applied aesthetic restorations may vary depending on the tooth's structural qualities (such as its pulp, enamel, and size), the child's cognitive and emotional state, and the material's cost (Srinath 2017; Pinkham JR et al. 2005). Crown restorations were first introduced in 1947 by the Rocky Mountain Company. In 1950, Engel introduced stainless steel crowns (SSC), but it was mostly through William Humphrey.

In 1971, a prominent opinion was ostensibly taken regarding the uses of stainless steel coin bail areas for deep bruises. If the vehicle of the 1970s is correct, it was put forward in the appearance of the ground protection of the SCCs. Webber introduced strip crowns in 1979. In 1983, Hartman developed the polished stainless steel crown for aesthetic anterior crowns. Buy Cheng crowns and Kinder crowns at the correct prices of the 1980s ("Available from: URL: <http://Www.Kinderkrowns.com/about-Us/History/> (Erişilme Tarihi: 30 Mart 2020)," n.d.). In the early 1990s, Randy replaced the band loop on the stainless steel crown by adding loops to the stainless steel crown. Again, the 1990s progressed. Norna Salon was made using a technical technique of dental restoration with a stainless steel crown without caries removal. Today, finding better systems and materials has brought zirconium to the forefront (Fellagh HF. 2016). Because of the high failure rate in restorations of teeth with excessive material loss, crown restoration is more preferred.

We can list the indications for crown restoration as follows;

- In teeth that have undergone endodontic treatment,
- In the treatment of advanced interface caries,
- In the treatment of broken teeth,
- In the presence of multifaceted caries,
- In children treated under general anesthesia,
- In patients with bruxism,
- In the treatment of defects in the cervical regions,
- In children with high caries risk and a lack of cooperation against rotary instruments,
- It is used in teeth undergoing pulpotomy-pulpectomy treatment. Desired properties in ideal primary tooth crown restorations;
- It must be aesthetic,
- chewing function should be preserved,
- The service life should be close to the life of the milk tooth,
- Protected teeth against breakage,
- The antagonist should not cause any wear on the tooth,
- It should be easy to apply,
- It should restore the lost function (Veerakumar R, Pavithra J, and Sekar K, 2017; Fellagh HF, 2016).

Today, the most commonly used crown restorations in the treatment of primary teeth are;

- Stainless steel crowns
- Polycarbonate crowns
- Strip crowns
- Pediatric jacket crowns
- Crowns created with the CAD/CAM system
- Zirconium crowns

Stainless steel crowns:

Restorative treatment may not always give the right response in primary teeth that have been severely damaged by caries. For this reason, in the early 1950s, Dr. Humphrey started to use stainless steel crowns for the first time as placeholders and crown restorations (Humphrey WP 1950). Thus, the frequency of indications for tooth extraction in children decreased, and the teeth began to stay in the mouth for a longer time. The stainless steel crown (SCC) contains 12–30% chromium, 8% nickel, and 1–20% carbon. It is reported that the lifespan of SCCs is longer than amalgams (Einwag and Dünninger 1996). Prefabricated SCCs have been used in the treatment of primary molars since 1950 (Engel R. J. 1950).

Over time, it has become more suitable for the anatomy of primary teeth and has been brought closer to the material structure that will facilitate applications (Ash, M. 1993). SCCs with traditional or veneered aesthetic surfaces are produced in many different sizes, can be corrected, stretched, and shaped with a bur, so they adapt well to crown margins, adhere well to tooth structures, and can be easily adapted to undercut areas. Due to these features, it stands out as a successful restoration type (Full, Walker, and Pinkham 1974; Duggal MS, Curzon MEJ, Fayle SA, Polard MA, and Robertson AJ 1995). It has been stated that SCCs are also produced in sizes suitable for first permanent molars so that they can be applied to permanent teeth. It is possible to shape and adapt these crowns to the teeth, and this is necessary for the success of the treatment. Restorations made with SCC have been considered a good alternative to traditional restorative treatments for many years, considering their durability and prognosis (N. P. T. Innes, Ricketts, and Evans 2007). It was stated by the British Association of Pediatric Dentistry in 2008 that restorations with SCC could be preferred (Kindelan et al. 2008). In a systematic analysis published by Innes et al. in 2015, evidence was revealed that SCC is the most appropriate restoration method, the effect of which is compared with traditional methods (Nicola P. T. Innes et al. 2015).

The indications for the stainless steel crown are as follows;

- Teeth with a small amount of enamel where adhesive bonding is not possible,
- Caries formation is seen on more than one surface of the tooth,
- Patients with poor oral hygiene,
- Root canal treatment and amputation were applied to milk teeth,
- In the presence of dental anomalies,
- In fractures with excessive substance loss that can be seen in primary and permanent teeth,
- They can be applied in subgingival caries (Einwag and Dünninger 1996; Seale 2002; Tayfun Alaçam, Uzel İ, and Alaçam A 2000).

In a study, Atieh compared the clinical success and retention time of PSC applied to primary molars after pulp treatment and the modified open sandwich technique using resin modified glass

ionomer cement and composite resin. In the study, which included 87 children aged between 4 and 7 years, 6, 12, 18, and 24 month controls were performed on 160 restorations. After two years of follow-up, the restoration rates were 95% for PSC and 92.5% for modified open sandwich restoration.

There was no significant difference between the two restorations in terms of marginal fit, proximal contact, occlusion, and recurrent caries criteria. When examined in terms of gingival health, better results were obtained in restorations made with the modified open sandwich technique compared to SCC (Atieh 2008). Webber observed some gingival enlargement in the follow-up of stainless steel crown restorations in a group of patients aged 8–12 years. However, it has been reported that gingival differences may be physiological in the mixed dentition and that the duration of use of the crown does not have a significant effect on gingival tissues (Webber 1974). Sharaf and Farsi, on the other hand, stated that the lack of interface contact by the SCC has no effect on gingival health. They reported that the most obvious factor for gingival health around stainless steel crowns is the level of oral hygiene (Sharaf and Farsi 2004).

Papathanasiou et al. evaluated the clinical success of different restorative materials. In this study, in which 604 restorations were examined in total, it was stated that the residence times of the applied restorations were SCC, amalgam, composite resin, and glass ionomer cement, respectively, from high to low. It has been reported that the mean residence time of SCC and amalgam restorations is 5 years. In this period, the percentage of PSCs remaining in the mouth is 68%, and amalgam restoration is 60%. After four years, the success rate of composite resin restorations is 40% and the average residence time in the mouth is 32 months; it has been reported that the success rate of glass ionomer cement restorations is 5% and the average residence time in the mouth is 12 months (Papathanasiou, Curzon, and Fairpo 1994).

There are studies that indicate that stainless steel crowns (which will be erased as their disadvantages) have a gingival compliance problem or (by looking at their condition inside the mouth) may cause periodontal problems. Tooth loss can also be seen in cases where the physician's clinical experience is insufficient or excessive cuts are made in the tooth. The aesthetic feature of stainless steel crowns is an important disadvantage not only for the anterior deciduous teeth, but also for the posterior deciduous teeth. In order to eliminate this negative feature, open-faced stainless steel crowns and veneered stainless steel crowns have been developed (Kindelan et al. 2008).

Although its durability, long life, and prevention of caries reoccurrence are the biggest advantages of SCC, studies have shown that the aesthetic expectations of patients and their parents cannot be met (Diana Ram, Fuks, and Eidelman, 2003; Townsend et al., 2014; Pani et al., 2016). Since these restorations did not meet the expectations in terms of aesthetics, various materials such as resin composite were tried to cover the buccal and occlusal surfaces, but these studies were not successful in clinical terms (Diana Ram, Fuks, and Eidelman 2003). In a study, it was stated that the aesthetics of dental treatments applied are one of the most important factors for parents (Zimmerman et al. 2009). For this purpose, single-faced crowns, strip crowns, and SCC veneer crowns have been developed (William F. Waggoner, 2002; 2015). Although these new materials contributed aesthetically, they caused some gingival problems (MacLean et al. 2007).

Open-face stainless steel crowns:

Although the aesthetic properties are insufficient and not accepted by parents and children, SCCs have continued to be used to protect the existing teeth in the mouth. With the advancement of technology, the aesthetics of stainless steel crowns have been improved. Open-faced stainless steel crowns are obtained by removing the front face of the stainless steel crowns we apply in the clinic and placing resin on the removed part. It was first used in the early 1980s (Wiedenfeld, Draughn, and Welford, 1994). In the survey study conducted by Nagarathna et al., 70% of the

participating physicians stated that they could make an open-faced stainless steel crown for the anterior region when faced with polyhedral caries. In addition, physicians stated that the ease of application is an advantage, and the child-parents are satisfied with this situation. 81.8% of the children showed a positive interaction during the treatment (Thimmegowda 2018). Disadvantages of open-faced stainless steel crowns; It is stated that the treatment period is long, fractures appear due to occlusal forces, and they are not aesthetically sufficient (Wiedenfeld, K. R., Draughn, R. A., & Goltra, S. E. 1995).

Veneered stainless steel crowns:

Due to the problems encountered with open-faced stainless steel crowns, veneered stainless steel crowns have been developed as an alternative. Veneered stainless steel crowns are a type of crown with an aesthetic appearance that can be cemented in the patient's mouth without any preparation. Initially, it was started to be used in the anterior region of deciduous teeth, and later it was applied to the posterior region of deciduous teeth as well. These crowns are applied in a single appointment and are reported to be less affected by the salivary environment. Since the aesthetic sections are made by the manufacturer, the time spent by the physician on the patient has also decreased (W. F. Waggoner and Cohen 1995).

Disadvantages of veneered stainless steel crowns;

- Contouring is performed only on the palatal or lingual surface, as there may be fractures on the facial surfaces during contouring,
- Color options are limited, so it is difficult to capture natural images,
- It is difficult to apply to children with limited space due to the volume's width.
- More teeth are cut due to passive seating in the tooth structure (W. F. Waggoner and Cohen, 1995; Roberts, Lee, and Wright, 2001) .

Fuks et al. examined the clinical success of veneered stainless steel crowns versus conventional stainless steel crowns over a 6-month period. At the end of the study, it was reported that the crown edge was compatible, the crowns were in the proper position/occlusion, proximal contact was provided, no bone resorption was observed, and no breakage or deformation was observed in the aesthetic part of the veneered stainless steel crowns. They stated that there was a significantly different gingival incompatibility in veneered stainless steel crowns compared to stainless steel crowns (Fuks, Ram, and Eidelman 1999).

Shah et al. examined 46 veneered SCCs they applied in the anterior region of 12 pediatric patients. They stated that the average duration of the crowned teeth in the mouth is 17.5 months (between 5 and 38 months). They reported that 24% of the crowns did not have resin fractures, and 61% of them did not have any fractures. When the gingiva of the crowned teeth were examined, they reported that 61% of the teeth had good gingival health, 24% had redness in the gingival area, and 15% had spontaneous bleeding (Shah, P. V., Lee, J. Y., & Wright, J. T. 2004).

Polycarbonate crowns:

Polycarbonate crowns, which have been widely used since the 1970s, are produced prefabricated as a result of the acrylic resin hardening with heat. As a result of the fabrication process, it is produced thinner than self-curing acrylic resin. Although aesthetically better than stainless steel crowns, they have lost popularity due to their fragility and ease of wear (Stewart, Luke, and Pike 1974). While applying the polycarbonate crown, a prefabricated polycarbonate crown suitable for the prepared tooth is selected and adhered to the tooth with acrylic resin after alignment. Restoration is completed by correcting the margins and polishing. It should be noted that polycarbonate crowns may break and completely separate from the tooth. Due to this insufficiency, their use is very low (Stewart, Luke, and Pike 1974) . Although the indications for

polycarbonate crowns are similar to those for stainless steel crowns, there are situations where they are contraindicated.

We can list them as follows:

- In cases with a deep bite,
- In the presence of progressive resorption in the root,
- In cases of excessive tooth wear or bruxism,
- In teeth with excessive substance loss,
- In cases of periodontal disease,
- In cases where a cross bite is seen,
- It is contraindicated in cases of deep bite (Venkataraghavan, Chan, and Karthik 2014; Nitkin, Rosenberg, and Yaari 1977).

Strip crowns :

Strip crowns have been frequently used in the restoration of primary teeth since the late 1970s. With the development of bonding agents and the increase in the bonding capacity of composites to enamel-dentin, strip crowns have been developed for the treatment of deciduous teeth in the anterior-posterior region in order to achieve tooth form, function, and aesthetics (William F. Waggoner 2002). Especially in anterior group milk teeth with excessive substance loss, strip crowns, which are much better aesthetically and have a wide usage area, are preferred instead of the stainless steel crowns that have been used for years. As an alternative to stainless steel crowns, strip crowns have been produced for posterior teeth, but they have not been used as widely as stainless steel crowns. Strip crowns are based on the selection of the appropriate mold in the form of plastic teeth for the tooth to be applied, the adaptation of the crown after the caries is removed, and the restoration of the entire tooth with a tooth-colored restoration material. After proper shaping, the resin material is cured with a light device. After that, the plastic tooth form is removed, and the remaining part of the tooth is corrected and polished (Citron 1995). The advantages of a strip crown are that it has good aesthetic properties, a shiny and smooth surface can be obtained, it is economical, a result that is very similar to the natural image can be obtained, and it can be applied easily and quickly (Kupietzky 2002). Tate et al., in a retrospective study investigating the restoration success of primary teeth treated with general anesthesia, examined the records of 504 patients and found PSCs to be significantly more successful than restorative methods such as amalgam, composite, and strip crowns. While the failure seen in SCCs was 8%, it was reported as 51% in strip crowns (Tate et al. 2002).

The main disadvantages are that the application technique is sensitive, blood and saliva contamination reduces the success of the restoration, sufficient dental tissue is needed for proper adhesion, and there are difficulties in its application because it requires cooperation in young children (Diana Ram, Fuks, and Eidelman 2003).

A sensitive technique is required during the application of strip crowns. In order to give the material a good shape and a suitable color, it is important to prevent its contamination with blood and saliva. In order for the bonding material and the composite to bond properly, the tooth tissue remaining after the preparation must be in sufficient quantity. At the same time, it can be difficult to apply strip crowns to very young children due to cooperation problems (Kupietzky 2002; D. Ram and Peretz 2000).

Pediatric jaket crowns:

It is an alternative type of crown used after severe caries or trauma. They are copolyester-containing, flexible, and naturally tooth-colored crowns. They can be shaped by cutting with

scissors. Thanks to the fact that some of the preparations can be made outside the mouth, the reduction of intraoral working time provides a significant advantage during its use, especially in children with a lack of cooperation. They are placed on the prepared tooth by applying acid and a bonding agent. If isolation can be achieved, either with composite resin cement or, if not, with resin modified glass ionomer cement, the jaket crown is adapted to the tooth and cementation is performed. The disadvantages of the pediatric jaket crown include wear by occlusal forces and discoloration over time. The cause of failure is usually the separation of the crown from the cement. When the copolyester crown is separated, the composite or resin-modified glass ionomer cement applied as an adhesive remains on the tooth and appears as a crown without the need for restoration (Daniels, Sim, and Simon 1966).

Crowns created with CAD/CAM:

In the last 25 years, CAD-CAM has become increasingly popular in dentistry. This technology enables the production of restorations with sufficient durability and a natural appearance with an application process that is easier, faster, and reduces the risk of human error (Zaruba and Mehl 2017).

Many restorative materials have been used to occlude primary teeth with excessive material losses, but a material that meets all expectations in terms of both durability and aesthetics has not been found. Due to the disadvantages of existing crowns, the need to produce crowns that can be prepared and applied more practically in parallel with technological developments and that have a much better aesthetic appearance has emerged. Today, the use of crowns prepared based on computer-aided design and production has come to the fore. The basis of this system; collection of data in a computer environment, the creation of a three-dimensional model, the realization of designs in the light of these data, and production using the realized designs. Computer-assisted shaping/computer-assisted restorations are called Computer-Assisted Design and Computer-Assisted Manufacture (CAD/CAM). CAD/CAM technologies were introduced to the world of dentistry in the 1980s. Since then, these technologies have continued to progress. Today, CAD/CAM systems cover indication areas such as inlays, onlays, laminate veneers, partial crowns, full crowns, and bridge systems. Although CAD/CAM restorations have begun to be used in pediatric dentistry clinical practice, their use in the restoration of primary teeth is not common, except for research (Ayşe METE 2014).

Inlay, onlay, laminate veneer crowns, bridges, skeletal structures of removable partial dentures, abutment in implant-supported prostheses, and infrastructure design and production in crown-bridge and hybrid prostheses constitute the indication areas of CAD-CAM systems. Materials currently used in CAD-CAM systems; ceramic (alumina, zirconia, and porcelain based) blocks, metal, and composite resin. Titanium and noble metals are used as restoration infrastructure (Willer, Rossbach, and Weber 1998; Piwowarczyk et al. 2005). Advantages of CAD-CAM systems; shortening the treatment time by reducing the number of sessions, eliminating the need for temporary crown preparation, reducing the rate of error, allowing the use of high quality materials, editing different parameters (such as restoration thickness, cement spacing) by the user, and archiving patient and restoration data (Miyazaki et al., 2009; Christensen, 2001). The disadvantages of CAD-CAM systems are the high production costs, the need for equipment and experienced personnel, the sometimes poor aesthetics of monochromatic blocks, and the difficulty of transferring to the computer environment in teeth with deep subgingival margin extension (Christensen 2001; Duret and Preston 1991). The fact that intraoral scanners in CAD-CAM systems have increasingly smaller designs allows their use in pediatric patients (Zaruba and Mehl 2017). CAD/CAM restorations are known to have become a common treatment for permanent teeth in children, but there are limited case reports of their use in primary teeth (Stines 2008).

CAD/CAM technology, restoration, and protective tooth preparation concepts should be considered together. Thus, unnecessary removal of the hard tissue of the tooth is avoided, and

restorations that are much more compatible and better processed can be obtained in a single session and are specific to the tooth to be restored. (Tsitrou and van Noort 2008) In a study in which the crown prepared by the CAD/CAM method in primary molars was compared in vitro, if there is extensive coronal destruction in primary molars, full coronal restorations prepared with the Chairside Economical Restorations of Esthetic Ceramic 3 (CEREC) system using polymeric CAD/CAM resin blocks are preferred because they are both aesthetically pleasing, have low preparation costs, and can be prepared at the bedside and in a single session. It has been reported that it can be done (Ayşe METE 2014).

Pediatric Zirconium Crowns:

Zirconium, symbol Zr and atomic weight 40, is a chemical element first discovered in Sri Lanka in the 18th century. The minerals of zirconium are zirconium silicate and zirconium oxide. Zirconium silicate is also called zircon, and zirconium oxide is also called zirconia. It has been reported that zirconium is not seen in nature in its pure form, but can be found in many different compounds. Zirconium is a highly resistant metal against temperature, wear, and corrosion. The purity of zirconium metal increases its mechanical properties. Since zirconium is a reactive metal, an oxide layer forms on its surface when it comes into contact with air or a solution. The oxide layer formed provides the zirconium with resistance to corrosion. It has been observed that the physical properties of zirconium are much superior to those of other ceramics. It is known that zirconium started to be used as a biomaterial in 1969 with the construction of hip prostheses by Hellmer and Driskell (Piconi and Maccauro 1999). Today, it is reported that stainless steel crowns are the most durable restorations that can be used in the treatment of deciduous molars and can protect the tooth structure the most. Despite the many positive features of SCC, researchers continue to develop new materials due to its poor aesthetics. Recently, zirconium crowns have been produced in order to eliminate the deficiencies in the restoration of milk teeth and to meet the aesthetic needs of children and parents. These materials can be used in the treatment of both milk incisors and milk molars. Prefabricated zirconium crowns were first introduced in 2008 under the brand name EZ-Pedo™ (Loomis, California, USA). The metal reflection from the gingiva, color mismatch, the presence of an artificial appearance in the stainless steel crown, and the absence of a personalized crown that can meet the aesthetic expectations of adults have led to the desire to produce a new material. With a patented system called Zir-Lock retention, grooves that provide mechanical retention were opened in the crown and they aimed to increase clinical success. They produced narrowed crowns for first and second primary molars whose mesio-distal size was reduced due to caries. Later, pediatric zirconium crowns with various features were produced by companies such as BruxZir (Glidewell), Lava™ Premium (3M ESPE; Katana ML; Kuraray), and NuSmile (Huston, TX, USA) (Prashant Babaji 2015). Tetragonal zirconium oxide polycrystals (yttria-stabilized tetragonal zirconia polycrystal: Y-TZP), stabilized by yttrium oxide, pass from one crystal phase to another, forming a resistant structure that stops crack formation and prevents crack propagation. Due to the high fracture resistance of the zirconium material, zirconium crowns are quite strong (Denry and Holloway 2010). The ready-made form of zirconium crowns released for children has been developed to be much stronger and more durable than the enamel structure. The transparency of zirconium crowns, thanks to their "translucent" feature, allows for the solution of the color problem that appears in the teeth after very good aesthetics and pulpal treatment (Tote et al. 2015). Holsinger et al., in a study in which they examined the clinical success and parental satisfaction of pediatric zirconium crowns used in anterior teeth, evaluated them according to their retention, gingival effect, color, contour, edge harmony, and abrasion on the opposite tooth. In the controls made as a result of an average of 21 months of follow-up, it was observed that no loss of retention was observed in any of the crowns, and 96% of them were present in the mouth. It was stated that no inflammation was observed in the gingival index (96%), no marginal alignment problem was observed in 86%, the color difference with the adjacent tooth (64%), was unnoticeable, secondary caries did not occur in the tooth with the zirconium crown,

and the opposing tooth. They reported no wear and tear. Parents also reported that the tooth with pediatric zirconium crowns had a natural tooth appearance (89%); it has been determined that children and parents are visually satisfied (Holsinger et al. 2016). In their case report, Cazaux et al. applied EZ-Pedo™/EZCrown™ pediatric zirconium crowns to the first primary molar after pulpotomy and followed up until the tooth exfoliated (29 months). They reported that 3 days after cementation, the gingival tissues regained their former health and appearance, there was no wear on the opposing tooth during the follow-up, and the tooth with a pediatric zirconium crown exfoliated at the same time (2 weeks apart) as the symmetrical tooth.

However, the researchers emphasized that it is often not easy to apply new materials in clinical practice, the application time of pediatric zirconium crowns is longer than that of stainless steel crowns in the first use, and the procedure becomes much faster and easier as clinical practice is gained over time. They reported that the application of the pediatric zirconium crown to the lower primary molars is easier than to the anterior teeth or upper primary molars. It is also emphasized that patient cooperation should be appropriate and that sedation may help crown application in uncooperative patients (Lopez Cazaux et al. 2017).

CONCLUSION:

Various treatment methods have been developed in cases of excessive substance loss in the teeth of pediatric patients. Factors such as the age of the child, aesthetic and functional needs, cost, ease of application, duration of treatment, and the number of teeth that require restoration constitute important criteria in determining the treatment methods to be applied. While aesthetics is in the foreground of a treatment method to be applied in the anterior region, function comes to the fore in the posterior region. There are not many randomized clinical studies on pediatric crowns.

According to the results of the existing studies, when stainless steel crowns and zirconium crowns are compared, it is known that patient satisfaction is higher with zirconium crowns, but due to the fact that the cost comes to the fore, stainless steel crowns and their more aesthetic modifications continue to be used today.

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A stomachache experience

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1-Introduction

Mesenteric ischemia is a condition that causes necrosis in the intestines due to insufficient oxygenation of the small and large intestines as a result of deterioration in mesenteric vascular blood flow. The most common condition of cronic mesenteric ischemia (CMI) is atherosclerosis in the splannic arteries, especially in advanced age (Jaster, 2016). In addition to atherosclerosis, vasculitides, fibromuscular dysplasia, radiation, aneurysms, arteriovenous fistulas, aortic dissection or coarctation, and congenital anomalies of the splannic vessels may also cause CMI. In diagnosis, doppler ultrasonography is being used with increasing frequency in the curvature of ripples in the deep and superior mesenteric artery (SMA). Angiography still continues to be the most sensitive method (Mastoraki, 2016). Although severe stenosis or complete obstruction of at least two of the three main splannic arteries is usually present due to the development of collateral segments, it may not have arisen in the classroom. Acute mesenteric vein thrombosis and consequently bowel infarction is one of the most feared cardiac controls. This kind of ability may not be easy to develop. Because, like many patients, there may be limitations due to common collaterals. Surgical revascularization and percutaneous transluminal angioplasty can be performed in two of the 3 splanchnic vessels in the mesenteric angiography in a patient who does not have any other gastrointestinal system disease and usually has symptoms of abdominal angina (Foley, 2016; Barret, 2015). A stent can be inserted during angioplasty. Bypass graft application to patients with S controls CMI is expressed as a safe treatment option (Flis, 2016). The average early operative mortality rate in elective revascularization is around 5% (0-12%). 80% of patients survive at 3-year follow-up. In the 3 years after revascularization, 15% of patients have s-graft features. Endo vessels have shelters in preferred places (Moghadamyeghaneh, 2015).

2-Case Report

1.st application

51 year old male patient; presents to the emergency department with the complaint of abdominal pain. Abdominal examination of the patient was unremarkable, mild leukocytosis (wbc: 12300) in the complete blood count and hematuria in the complete urinalysis, the urinary system was evaluated as normal in the abdominal usg. In the right lower quadrant of the abdomen, at the level of the terminal ileum and cecum, increased thickness of the intestinal wall, increased peristalsis, and fluid distension were observed, and abdominal CT is performed upon the recommendation of the radiologist. In the General Surgery consultation, it was learned that the

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abdominal pain and severity of the patient that started after the meal increased. The patient did not have a cardiac disease, has a one pack/day smoking history and a history of substance abuse. Due to widespread tenderness and rebound in the abdomen, the patient was admitted to the general surgery clinic with the diagnosis of acute abdomen, and preoperative preparations and resuscitation were started. Oral intake of the patient, whose ASA 2 anesthesia risk was determined, was stopped and fluid resuscitation, nasogastric decompression and antibiotherapy was started. The patient was discharged voluntarily without waiting for the results of the examinations. In the patient's entire abdominal contrast-enhanced CT examination; Fluid distention, dilatation and air fluid levels were observed in the ileal loops and were evaluated as compatible with ileus, while the increase in fringing-like density in the mesentery fat areas was recommended to be evaluated primarily in terms of inflammatory ileitis and terminal ileitis.

2.st application

Four days later, she came to the emergency room with the complaint of abdominal pain again. In general surgery consultation; There was pain and rebound in all quadrants of the abdomen. Contrast-enhanced tomography of the entire abdomen was requested, considering mesenteric vascular disease. Here, an exophytic localized irregular noncalcified plaque that causes more than 80% stenosis in the lumen of the superior mesenteric artery (approximately 2 cm segment at the exit of the SMA) was observed (Picture 1: stenosis in SMA). Thromboembolism causing total occlusion of the end branches of the SMA was observed in the pelvic region. All this Considering the imaging results and the patient's history, he was referred to another center for the treatment of stenosis in the SMA with interventional radiological methods for the treatment of abdominal pain due to intestinal hypoperfusion due to microemboli and narrowing of the SMA.

3.st application

A stent was placed in the SMA by endovascular method in an external center.

4.st application

The patient, who has applied to our clinic with the complaint of abdominal pain again 2 months later, states that this abdominal pain, which increases with meals, prevented him from eating anymore, he was getting weaker and he cannot stand these pains. Imaging examinations of the patient, who was hospitalized because of diffuse tenderness in the abdomen, showed a stent in the SMA and left common iliac artery (Picture 2: CT view of the stents). In the left lower quadrant, thickening of the intestinal wall, distension and edema, and minimal free fluid between the intestinal loops were observed. Oral intake of the patient was stopped, the patient is relieved by the continuation of anticoagulant treatment, fluid resuscitation and the use of analgesics.

5.st application

One month after discharge, the patient, who was admitted to another center with complaints of abdominal pain and inability to pass gas and stool, has followed up with the diagnosis of ileus. The abdomen distended and tenderness in the left lower quadrant and a palpable mass in the rectum were felt. Laboratory examinations were normal, imaging examination (whole abdomen with contrast CT) has cholelithiasis, cortical cyst in bilateral kidneys, collection area associated with the anterior wall of the middle part of the rectum, a stent was present in the proximal 4 cm segment of the SMA and its lumen was open, the lumen of the stent in the left common iliac artery. Occluded throughout, extending between the small intestines in the retrovesical space in the pelvis, continuing towards the pelvic wall on the left, the large 50*30 cm dimensions, which cannot be clearly distinguished from the neighboring intestines, were observed to be related to each other. The patient, who was followed up with antibiotherapy with the diagnosis of pararectal abscess, was discharged with anticoagulant treatment after his general condition improved and the size of the abscess decreased in the control.

6.st application

The diagnosis of acute abdomen was operated in an external center and small bowel perforation has been observed. Small bowel resection, ileostomy and mucous fistula have been performed.

7.st application

The patient was a cachectic appearance, insufficient oral intake, and fluid and electrolyte imbalance. He stated that he has lost 30 kg of weight in the last 6 months. Hyponatremia was treated.

3. Discussion

There are 3 arteries in the mesenteric system. The celiac artery is the superior mesenteric artery (SMA) and the inferior mesenteric artery (IMA). Celiac artery; proximal of stomach and duodenum, SMA; distal duodenum, jejunum, ileum, up to 2/3 of transverse colon, IMA; Starting from the 1/3 distal transverse colon, it supplies the rectum and proximal anal canal. There is rich collateral circulation between the branches of the mesenteric vessels. Therefore, clinical findings occur when 2 or 3 main arteries are occluded or have critical stenosis. Most commonly, the obstruction of the SMA, which is described as acute mesenteric ischemia, is encountered as a result of embolism or thrombosis developing on the basis of chronic atherosclerosis. In addition, when bowel ischemia is encountered without an occlusive lesion, non occlusive mesenteric ischemia is defined. Although it is thought that almost all ischemia originates from the arterial system, intestinal ischemia of venous origin can be seen. Acute mesenteric ischemia accounts for 1 out of 1000 hospitalized patients. It is a fatal disease if left untreated. Although early intervention provides a complete recovery, it is a difficult disease to diagnose (Tilsed,2016). It is a disease that should be kept in mind when the patient has recurrent episodes of abdominal pain that starts after meals and lasts for 1-3 hours, as in our case example of CMI, which has been admitted to the hospital 7 times and was followed up every time. Although the pain may be mild at first, it can become more severe in weeks and months. Postprandial vasodilation reduces peripheral resistance, but flow cannot increase in the presence of proximal fixed occlusive lesions, resulting in transient ischemic pain. This pain has been described as bowel angina. Weight loss may occur because patients are hesitant to eat. Nonspecific findings such as hypoalbuminemia, hypoproteinemia, and hypocholesterolemia caused by malnutrition may be encountered. Our case was cachectic at the 4th admission and did not want to take oral. In our patient, a stent was inserted during angioplasty with an endovascular treatment approach. There are publications stating that surgical or endovascular interventions have similar success rates (4, 7), but in some publications, bypass grafting is indicated as a safe treatment option in patients with symptomatic CMI(6).

4. Conclusion

Chronic mesenteric ischemia, which we do not encounter very often in our general surgery practice; Due to the difficulty of treatment, the patient progresses to death with intestinal ischemia at the end of the treatments and follow-ups for abdominal pain attacks and weight loss symptoms.

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Conflicts of interest

The authors declare that there are no potential conflicts of interest relevant to this article

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Picture1: stenosis in SMA



Picture 2: CT view of the stents

State-Trait Anxiety Levels Of Teachers During The Covid-19 Pandemic In Kırşehir, Turkey

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Introduction

Anxiety is a feeling of unease, which can be experienced by the individual in the face of events that are assumed to be dangerous or threatening without any apparent reason. Anxiety is an unpredictable, unpleasant, and overwhelming feeling, which is associated with sad feelings about the future or anticipation of something bad about to happen. It is a feeling of nervousness and stress because of generalized thoughts around bad things that could happen with no identifiable cause (Şahin, 2019). Anxiety disorders are described in DSM-V and categorized as different types such as generalized anxiety, panic, and social anxiety disorder, agoraphobia, specific phobia, and separation anxiety disorder (DSM-V, 2013). Everybody can feel anxious time to time. Anxiety is examined in two categories, which are the “state anxiety” and the “trait anxiety”. Trait anxiety is a generalized and persistent state of anxiety that is not tied to a specific event or situation. In contrast, state anxiety occurs before or during the so-called jeopardous situations. This is the anxiety experienced by almost every individual, which others can understand, and which can be attributed to logical reasons (Yıldız, 2010).

Infectious disease outbreaks induce anxiety and worry, causing individuals to feel ashamed, unsuccessful, and weak (Sim, 2010). Furthermore, psychological reactions occurring during outbreaks vary in a spectrum extending from extreme forms of anxiety and fear to feelings of indifference. The long-term psychological effects of outbreaks include the weakening of social and interpersonal ties. The psychological consequences of such periods are not always short-lived. Experiences in such periods can cause severe and persistent mental disorders as well (Huang & et al., 2020, Zhu & et al., 2020). Anxiety, mood and post-traumatic stress disorders are the most common mental distresses in epidemics (Tükel, 2020). Indeed, health anxiety experienced during outbreaks is an anticipated situation. People may develop a state of mental exhaustion due to uncontrollable health anxiety, which consequently can induce a vicious cycle. Another form of health anxiety during outbreaks is the anxiety of transmitting the disease to someone (Pierce, 2020). COVID-19 outbreak is the most recent one that emerged in China in 2019 and then spread around the world turning into a pandemic. When pneumonia cases of unknown origin increased in Wuhan (China), China reported this situation to the World Health Organization (WHO) on 31 December 2019. Chinese scientists named this coronavirus “2019-nCoV” on January 7, 2020. Spread around the world in a short time, the coronavirus started to have been identified in many people in Turkey as of March 10, 2020. From January 3, 2020 to January 10, 2022, there were 9,977,561 confirmed cases of COVID-19 in Turkey, with 83,702 deaths reported to WHO (WHO, COVID-19 Dashboard, 2022). The COVID-19 pandemic caught people unprepared and defenseless, affecting them globally and reaching alarming levels all over the world. The lives of many people were lost or disabilities occurred, leading to the development of psychological problems (Pektekin, 2020).

In the study of Nelson et al. (2020) in March 2020, it was revealed that 67.3% of 9009 participants were very much or extremely worried about COVID-19 (Nelson & et al., 2020). In the study of Lai et al. (2020) in China, 44.6% of healthcare professionals were found to experience anxiety (Lai & et al., 2020). Anxiety also arises from loneliness and limited social contacts during quarantine and isolation periods of outbreaks. People start experiencing anxiety because they cannot run their business and fail to meet the needs of their dependents (Brooks & et al., 2020, Huremović, 2019). Emotional responses to COVID-19 are experienced by all working individuals from all social levels, primarily healthcare professionals and people that are directly affected by the untoward consequences of the pandemic (Sim, 2010, Huang & et al., 2020, Zhu & et al., 2020, Stuijzand & et al., 2020, Pappas & et al., 2009)

During the COVID-19 pandemic, schools have spent their best efforts to carry out educational activities online. All educational activities were interrupted, affecting students, teachers, and parents remarkably. Literature review revealed that many studies reported that the COVID-19 pandemic induced many feelings affecting the mental status of students unfavorably, such as learning and exam anxiety, worry, fear, extreme levels of stress, irritability, aggression, insomnia (Kara, 2020, Sintema, 2020, Cao & et al., 2020, Daniel, 2020). However, the number of studies investigating the problems experienced by teachers during the COVID-19 pandemic is not adequate despite their key roles in education. Studies available in the literature are mostly about the views of teachers on education (Ozamiz-Etxebarria & et al., 2021, Wakui & et al., 2021, Wong & et al., 2021, Çakın & Akyavuz, 2020, Çiçek, Tanhan & Tanrıverdi, 2020, Li, 2020, Stachteas & Stachteas, 2020). It has been determined that more international studies are needed to determine the effect of the epidemic on teachers' mental states (Ozamiz-Etxebarria & et al., 2021). The teacher with a high level of anxiety cannot fully devote herself to her task. Excessive anxiety causes the individual to become incompetent and result in disagreeable and irritable behavior in the workplace. On the contrary, lack of anxiety causes the individual to behave inconsiderately. Therefore, the anxiety level of the teacher in the workplace should not be at extremes (Başar, 1999).

Cooperation across professional associations, including those of psychiatrists, nurses, psychologists, and social workers, plays a crucial role in the resolution of psychosocial problems occurring during the pandemic. In this context, the Online Mental Health Program for Coronavirus Infection (KORDEP) was implemented to provide psychosocial support to people affected by the COVID-19 pandemic in Turkey. This program is considered an important step in coping with difficulties occurring while we live the pandemic (Pektekin, 2020). With awareness about the anxiety levels of teachers, such psychological support programs may provide more effective guidance.

Based on these considerations, we designed this study to examine the anxiety levels of teachers during the COVID-19 pandemic.

Patients and Methods

The research was conducted on 1704 volunteering teachers working in primary, secondary, and high schools, and in institutions providing pre-school education in Kırşehir (located in central Anatolia) province in the 2020-2021 academic year. Teachers were reached by the snowball sampling method. With this method, teachers participating in the research were asked to distribute data collection tools to their colleagues.

Research data were collected using an online demographics form, the State Anxiety Inventory (SAI) (STAI FORM TX-I), and the Trait Anxiety Inventory (TAI) (STAI FORM TX-II). The design and adaptation of the inventories into Turkish was performed by Öner and Le Compte (Öner & Compte, 1983). Both inventories comprise 20 items to be scored using a 4-point Likert

scale. Each item can be rated from 1 to 4. Response options in SAI were as follows: (1) not at all, (2) somewhat, (3) moderately so, (4) very much so. Response options in TAI were as follows: (1) almost never, (2) sometimes, (3) often, (4) almost always. The inventories included direct and reversed items or items phrased in the semantically opposite direction. The reversed ones were the items 1, 2, 5, 8, 10, 11, 15, 16, 19, and 20 in SAI. In TAI, the reversed ones were the items 21, 26, 27, 30, 33, 36, and 39. The minimum and maximum scores on both inventories could be 20 and 80, respectively. A high score indicates high anxiety, a small score indicates low anxiety. In this study, Cronbach's alpha values of the inventories were calculated as 0.951 for SAI and 0.891 for TAI. It has been reported that a scale with an alpha coefficient of 0.80-1.0 is highly reliable (Alpar, 2011). According to the calculated alpha coefficients for this study, our results are reliable.

Data collection forms were delivered to teachers online. Before teachers started filling out the online data collection forms, the aim of the research was explained to them in accordance with the Declaration of Helsinki and consent was obtained from participants. It took about 10-15 minutes for teachers to fill out the forms.

The data of the study were evaluated in computer environment. Kolmogorov-Smirnov, Shapiro-Wilk, Levene test, percentage, mean, standard deviation, independent t test, ANOVA, Duncan multiple comparison test and Cronbach's alpha coefficient were used in the analyzes.

Results

Of the teachers, 47.1% were women and 52.9% were men. Descriptive statistics of demographics and comparative answers to inventory items are presented in Table 1. As seen in Table 1, the effects of the gender variable on both SAI and TAI scores were statistically significant ($p < .001$). Women's state and trait anxiety scores were higher than those of men. The number of children of a participant affected the state anxiety level ($p < .05$). The state anxiety levels of those with only one child were higher than those of the rest of the participants. The effect of the number of children had a statistically insignificant effect on the trait anxiety level ($p > .05$). The participants' marital status, education level, length of service, and the type and location of the school of employment did not affect their state and trait anxiety levels ($p > .05$). The effect of job tasks performed by the participants at the time of the study on the state and trait anxiety levels was statistically significant ($p < .05$ and $p < .001$, respectively). The state and trait anxiety levels of participants working as administrators were lower compared to those performing other jobs.

Table 1. Descriptive characteristics and comparison of scale scores (N=1704)

Demographics	characteristics	n (%)	Mean±SD	p
Gender				
State Anxiety	Men	802 (47.1)	39.17±11.46	0.000
	Women	902 (52.9)	41.93±11.04	
Trait Anxiety	Men	802 (47.1)	40.28±7.70	0.000
	Women	902 (52.9)	43.38±7.71	
Marital Status				
State Anxiety	Single	163 (9.6)	41.49±11.60	0.307
	Married	1541 (90.4)	40.54±11.29	
Trait Anxiety	Single	163 (9.6)	43.01±8.95	0.063
	Married	1541 (90.4)	41.80±7.72	
Number of children				
	0	179 (10.53)	40.71±10.42 ^a	0.017
	1	377 (22.18)	41.31±11.52 ^a	
State Anxiety	2	898 (52.85)	40.94±11.58 ^a	
	3	191 (11.24)	38.67±10.48 ^{ab}	
	4+	54 (3.17)	37.44±10.81 ^b	
Trait Anxiety	0	179 (10.53)	42.24±8.68	0.185
	1	377 (22.18)	42.39±8.08	

	2	898 (52.85)	41.94±7.64	
	3	191 (11.24)	40.94±7.32	
	4+	54 (3.17)	40.57±8.45	
Educational status				
State Anxiety	Bachelor	1532 (89.9)	40.54±11.37	0.347
	Postgraduate	168 (9.9)	41.41±10.86	
Trait Anxiety	Bachelor	1532 (89.9)	41.92±11.37	0.936
	Postgraduate	168 (9.9)	41.97±7.13	
Length of service				
	1-5 years	114 (6.7)	39.15±11.71	
State Anxiety	6-15 years	711 (41.7)	41.17±10.71	0.151
	16-25 years	614 (36.0)	40.68±11.56	
	26+ years	265 (15.6)	39.71±12.10	
Trait Anxiety	1-5	114 (6.7)	42.44±8.87	0.054
	6-15	711 (41.7)	42.50±7.74	
	16-25	614 (36.0)	41.44±7.75	
	26+	265 (15.6)	41.23±7.86	
Type of school of employment				
	Pre-school	89 (5.2)	41.32±11.17	
State Anxiety	Primary school	453 (26.6)	40.92±11.29	0.722
	Secondary school	576 (33.8)	40.10±11.12	
	High school	509 (29.9)	40.79±11.60	
	Other	77 (4.5)	41.0±11.41	
Trait Anxiety	Pre-school	89 (5.2)	42.13±8.16	0.796
	Primary school	453 (26.6)	42.07±7.74	
	Secondary school	576 (33.8)	41.86±7.92	
	High school	509 (29.9)	41.69±7.78	
	Other	77 (4.5)	42.80±8.24	
Location of school	Village	85 (5.0)	41.02±11.21	
	Town	46 (2.7)	43.19±10.68	
State Anxiety	District	440 (25.8)	40.80±11.17	0.406
	City center	1133 (66.5)	40.43±11.41	
	Village	85 (5.0)	43.05±8.69	
Trait Anxiety	Town	46 (2.7)	42.95±8.28	0.428
	District	440 (25.8)	41.84±8.16	
	City center	1133 (66.5)	41.83±7.65	
Job title	Teacher	1486 (87.2)	40.86±11.31	
State Anxiety	Administrator teacher	169 (9.9)	38.28±10.82	0.015
	Other	49 (2.9)	41.73±12.31	
Trait Anxiety	Teacher	1486 (87.2)	42.17±7.75	0.000
	Administrator teacher	169 (9.9)	39.57±8.18	
	Other	49 (2.9)	42.38±8.58	

*Groups with significant differences in between, as identified by Duncan's multiple comparisons, are marked with different letters in the same column.

Descriptive statistics of 40 items in both inventories used in the study are presented in Table 2. The mean scores of participants on SAI and TAI were 40.6(11.3) and 41.9(7.8), respectively. The state-trait anxiety levels of teachers were moderate.

Table 2. Mean scores from the State-Trait Anxiety Inventory

Scale	Mean
State Anxiety Inventory	40.63±11.32 (min= 20, max= 80)
Trait Anxiety Inventory	41.92±7.85 (min=20, max=71)

Discussion

In this study, teachers' anxiety levels were found to be of moderate severity. Studies in the literature support this finding. According to the results of a study conducted during the SARS epidemic in Hong Kong, people were found to experience moderate levels of anxiety (Leung & et al., 2003). It was previously reported that teachers' anxiety about the COVID-19 pandemic was of moderate severity (Çiçek & et al., 2020). In the study performed by Wong et al. (2021), it was found that teachers experienced depression and stress, and their anxiety was mostly about themselves and their family members. In that study, it was found out that teachers also had anxiety about distance education (Wong & et al., 2021).

According to Fuller (1969), teachers' feelings of anxiety occur at certain time points normally (Fuller, 1969). Teachers experience anxiety when faced with certain concerns such as using time effectively in the classroom, completing activities effectively, and maintaining discipline in the classroom (Gangal, 2020). Teachers may also develop anxiety in response to some situations, such as incompetency in the field of the job task, lack of knowledge and experience, or the inability to use relevant knowledge and equipment (Uçak & Bindak, 2017). Nevertheless, teachers need to control the level of anxiety they experience in the workplace. It is a favorable finding that teachers' anxiety levels did not increase during the pandemic and remained at moderate levels. Teachers, who perform education and training activities online and remotely during the pandemic, may experience some difficulties in communication. The methods used by teachers for student motivation regarding education and the maintenance of good health are very important for the assessment and improvement of educational activities. It has been observed that teachers have readily adapted themselves to the use of distance education during the pandemic. The fact they could manage the process well might help prevent them from developing increased anxiety levels.

In this study, we found that the marital status, education levels, and lengths of service of teachers did not affect their state and trait anxiety levels. These findings are consistent with those reported in the literature (Çiçek & et al., 2020, Sakaoğlu & et al., 2020, Çifçi & Demir, 2020, Bulut & Duman, 2021). In a study performed by Sakaoğlu et al. (2020), the examination of anxiety levels by education status showed that there were no differences between the state and trait anxiety scores. Similar to the findings of the present study, it was also reported in the literature that the level of anxiety increases as the education level of teachers increases (Sakaoğlu & et al., 2020). In the study performed by Bulut and Duman (2021), no relationships were found between the length of service and the level of anxiety of participants (Bulut & Duman, 2021). Some studies reported higher anxiety levels in female, young, and single teachers compared to their male, older, and married counterparts during the COVID-19 pandemic (Çiçek & et al., 2020, Çifçi & Demir, 2020). The present study found that women's anxiety levels were higher than that of men. In studies conducted on different sample groups in the literature, it was found out that women were more worried than men during the COVID-19 pandemic (Bulut & Duman, 2021, Çölgeçen & Çölgeçen, 2020, Kayaoğlu & et al., 2021).

In the present study, the state anxiety levels of teachers with only one child are higher than those of their counterparts. Being a child without siblings in a family means receiving all focused attention and care. The low anxiety levels of parents with no children or more than one child suggest that attention directed to the only child in the family can be potentially excessive. Furthermore, such increased anxiety levels may result from the fear of the likelihood of leaving the only child all alone in case of conditions that could put the parents in peril.

Among the participants, the state and trait anxiety levels of teachers in administrative positions were lower compared to the rest. This may be due to the better command of the pandemic-related

subjects by the administrators, who acquire information through frequent meetings and by closely following up the state and government policies about the pandemic.

Following the start of the COVID-19 pandemic in the world and our country, telepsychiatry applications allowing telephone calls or online video calls were readily implemented, providing access to all healthcare professionals and anyone affected by the pandemic. This has led to the development of a mental support mechanism for individuals. Solidarity and charity work in society are critical in alleviating the pandemic-related problems(Tükel, 2020).

Conclusion

It is pleasing that teachers have not developed severe levels of anxiety and worry during the pandemic. This has helped prevent the serious interruptions in the education system. During periods of outbreaks, people should be in social, psychological and, when necessary, material solidarity. In such a process, people should do their parts.

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Intracalcaneal Lipoma

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Introduction

The lower extremity is one of the most common anatomical sites for bone and soft tissue tumors and tumor-like lesions. Primary bone tumors are rare, and most are benign lesions. Of these, osteochondroma, enchondroma, osteoid osteoma, chondroblastoma, giant cell tumor, chondromyxoid fibroma, liposclerosing myxofibroid tumor, and intraosseous lipoma are the most common lesions.

Intraosseous lipoma, among the primary benign bone tumors, is less commonly diagnosed than other primary bone tumors and may occur incidentally in radiological examinations performed for another reason if they are asymptomatic. Therefore, the true incidence cannot be determined clearly.

The primary and only reason for patients presenting with intracalcaneal lipoma to apply to the clinic is chronic heel pain that persists despite analgesic therapy and worsens with walking. In this benign bone tumor where spontaneous regression is possible, the most critical reason that leads the orthopedist to surgery is the patient's complaint of pain and, more importantly, the radiological resemblance of the lesion to other bone tumors. In particular, intracalcaneal lipomas can be radiologically mistaken for aneurysmal bone cysts, giant cell tumors, unicameral bone cysts, pseudocysts, and bone infarcts (Arkun, 2016; Gero & Kahn, 1988). In intraosseous lipomas, for which curettage and grafting are sufficient, no recurrence is observed after surgery (Levin et al., 1996).

Bone-related lipomas are localized in intraosseous, cortical, or parosteal zones and originate from lipocytes. The most common one is intraosseous localization. This rare lesion accounts for 0.1% of tumoral lesions of the bone (Hassani et al., 2014; Unni & Inwards, 2010). The sex predilection in intraosseous or intramedullary lipoma is controversial (Goto et al., 2002). Intraosseous lipoma was defined in three stages by Milgram et al. (Milgram, 1988a) according to the pathological changes in the internal structure of the lesion (Table 1). An intraosseous lipoma detected in Stage 3 can be mistaken for a bone infarction.

Table 1. Milgram Staging

Stage 1	There are sharply demarcated lipocytes of mature homogeneous adipose tissue replacing trabecular bone. A radiolucent lesion with fine marginal sclerosis and slightly expanded well-defined margins can be observed on plain radiographs.
Stage 2	Foci of necrosis and calcified areas may be determined in the lesion center of mature adipose tissue cells. At this stage, the cystic area accompanies the lesion.
Stage 3	There is increased fat necrosis, coarse calcifications, myxomatous degeneration of fat cells, and larger cystic areas. There is an increased density of calcification in an expanded, well-defined, radiolucent lesion on direct radiography. The sclerotic zone around the lesion is thicker.

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Case Presentation

A 52-year-old male patient, who was admitted to the outpatient clinic with right plantar heel pain, described the symptom as right heel pain that increased significantly after getting up in the morning and walking a few steps and got worse after a long-standing activity or lifting a heavy load. Although he could not remember when this pain started, it was recorded that the patient's complaints had increased in the last year, and there was no previous trauma to his right heel in the anamnesis.

On physical examination, there was pain on palpation, especially on the plantar face. No tenderness was detected on the medial and lateral face of the heel. Ankle and subtalar joint ranges of motion were complete both actively and passively. A preoperative visual pain scale was performed on the patient. The patient rated his pain with a score of 7 to 8 out of 10. Two-way radiography of the ankle was requested from the patient. On the radiograph, it was seen that there was a 25x30 mm unicameral expansile lesion in the calcaneal medulla, hypodense compared to the intratrabecular bone density of the calcaneus, and hyperdense ossification foci in the center of the lesion (Figure 1). Contrast-enhanced ankle magnetic resonance imaging (MRI) was performed for further examination. MRI images reported that a lesion compatible with lipoma was observed in the medulla, which was 36x20 mm in size in the calcaneus, isointense in well-defined T1W, suppressed in fat-suppressed sequences (Figure 2a,2b). After intravenous contrast agent injection, two cystic foci, the largest of which was 9x5 mm, were noted in the center of this lesion. Both MRI and plain X-ray findings favored Milgram grade-3 intraosseous lipoma. (Figure 2c,2d,2e,2f)

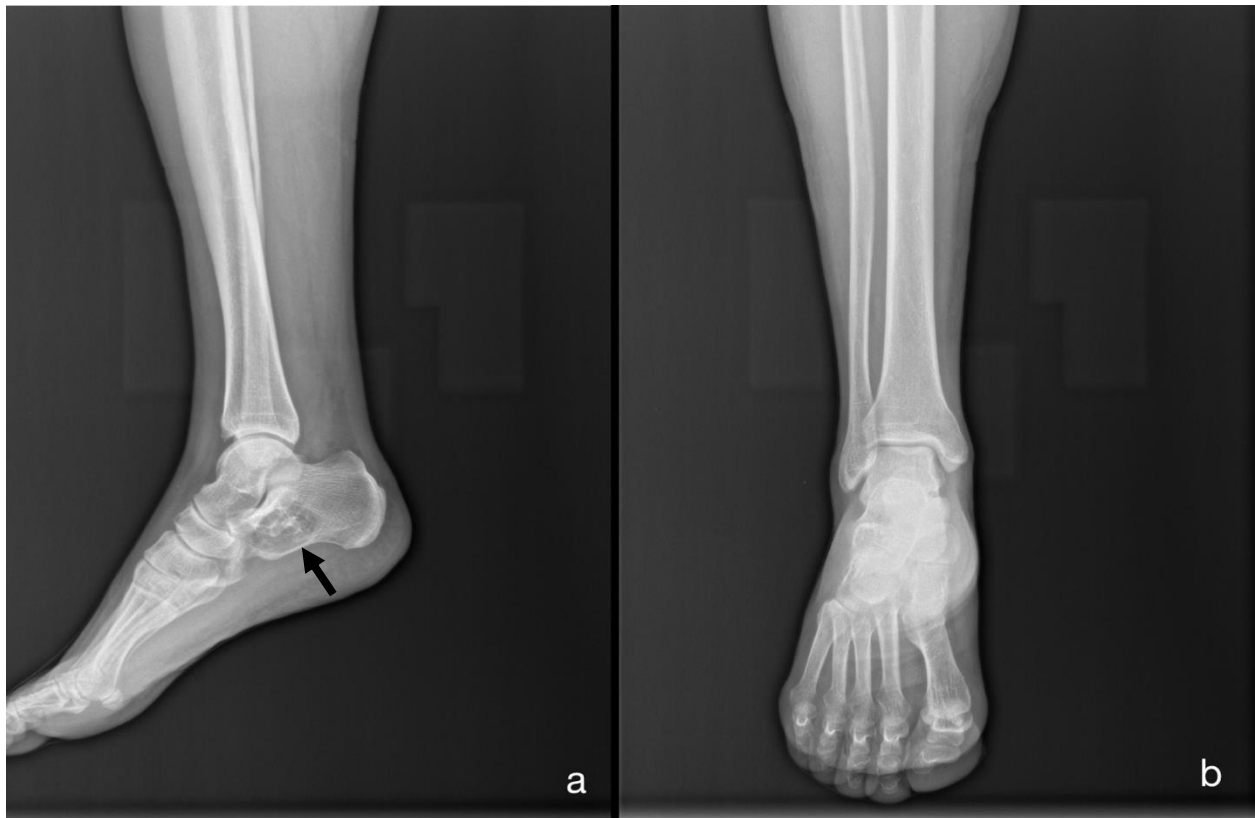


Figure 3 a: Cystic lesion is seen where the arrow points in lateral ankle X-Ray view, b: Anteroposterior (AP) ankle X-Ray view

As a result of the physical examination and tests, no infective or degenerative foot and ankle pathologies were considered in the patient, and the diagnosis of intracalcaneal lipoma was made. The patient was explained that the pain in this disease could be corrected surgically. The complications that may occur were explained to the patient by giving preoperative counseling services. Surgical preparation of the patient, whose informed consent was obtained, was planned.



Figure 4 Magnetic resonance (MR) imaging a: Sagittal T1 FSE Non-contrast (NC) MR view, b: Axial T1 FSE NC MR view, c: Sagittal T2 FSAT MR contrast view, d: Axial T2 FSAT MR contrast view, e: Sagittal T1 FSE MR contrast view, f: Axial T1 FSE MR contrast view

It was decided whether the patient would be treated with curettage grafting alone or plating in addition to this treatment based on Mirels' prophylactic detection criteria (Mirels, 1989). A locked calcaneal plate was applied as prophylactic fixation material to the patient who scored 9 according to this criterion. The patient was placed in the lateral decubitus position. The mass in the calcaneus was reached with an extended lateral approach. After the calcaneal cortex was removed, the fluid-filled yellow cystic structure was aspirated, and an excisional biopsy was performed to send the mass like fat globulins in the cyst to pathology. It was observed that the area around the evacuation area was sclerotic. The surrounding of this sclerotic wall was decorated with the help of a drill, and the connection of this evacuated area with the bone medulla was ensured. A bone graft was then placed in this area. To increase the efficiency of the graft, tricalcium phosphate, a synthetic ceramic graft with osteoconductive properties, and an autogenous bone marrow graft with osteoinductive properties

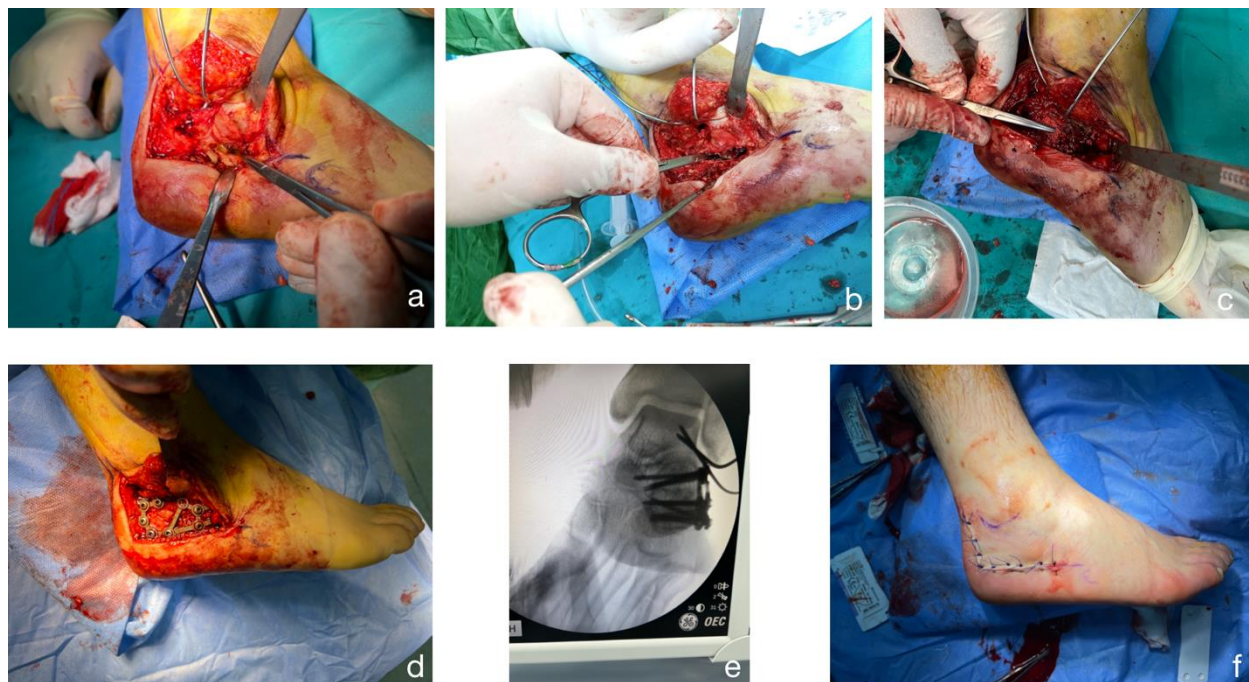


Figure 5 a: Intraoperative view of the cystic structure before opening in the calcaneus' cortex, b: Intraoperative view of the cystic structure after opening in the calcaneus' cortex, c: Curettage and graft placement of the cystic structure in the calcaneus, d: Applied view of locked calcaneal anatomical plate, e: Intraoperative fluoroscopic AP view after plate use f: Wound closure of extended lateral approach view

were combined. Then, a locked calcaneal anatomical plate was applied, and the operation was completed without complications. (Figure 3)

The patient was followed up with a splint for three weeks after the operation, and weight bearing was started approximately five weeks after the operation. Complete healing was determined radiologically seven weeks after the operation. (Figure 4) Clinically, the patient was followed up for 20 weeks after the operation. The patient's visual pain scale was measured. The patient's pain was recorded by giving a value between 2 and 3 points out of 10. As a result of the postoperative controls, it was decided that the bone graft area was healed, both by showing that bone remodalization was achieved on the radiograph and by evaluating the patient's clinic.



Figure 6 a: Early postoperative (EP) AP ankle X-Ray view, b: EP lateral ankle X-Ray view, c: 5 week after operation AP ankle X-Ray view, d: 5 week after operation oblique ankle X-Ray view, e: 7 week after operation AP ankle X-Ray view, f: 7 week after operation oblique ankle X-Ray view

Discussion

As reported in the literature, intraosseous lipomas develop in less than 0.1% of all bone tumors and have an incidence between 0.1% and 2.5% among primary bone tumors (Hassani et al., 2014; Unni & Inwards, 2010). The first case was reported in the femoral shaft by Brault in 1868 (Milgram, 1988b). The second case was published in the femur diaphysis in 1901, and the third was published in the fibula in 1910 (Unni & Dahlin, 1996). Intraosseous lipomas are most commonly located in the subtrochanteric area of the proximal femur, second most frequently in the calcaneus (Hong et al., 1999).

Intraosseous lipomas are asymptomatic and are often detected incidentally. As in our case, 70% of the cases in symptomatic patients present with pain, and it is thought that this pain is caused by micro-fractures in the trabecular network of the calcaneus due to axial loading (Goto et al., 2002).

When the literature is reviewed, neoplastic and non-neoplastic pathologies such as simple bone cysts, aneurysmal bone cysts, osteblastoma, enchondroma, chondromyxoid fibroma, non-ossifying fibroma, giant cell tumor, chondroblastoma, fibrous dysplasia, and chondrosarcoma are considered in the differential diagnosis (Futani et al., 2007). Pain is the primary surgical indication in these masses, and among the surgical options, curettage of the cystic area followed by grafting or cementing, if necessary, is preferred to fill the defect area. After filling the defect area, it was decided to use fixation material according to Mirels' prophylactic fixation criterion (Azarsina et al., 2019) in some studies against the risk of fracture of this bone area. In contrast, in some studies, it was decided to detect the mass after the curettage of the mass with the concept of a critically sized cyst (Pogoda et al., 2004).

In this case, the patient was evaluated as Milgram stage 3. The use of fixation material was evaluated according to Mirels criteria. According to this criterion, getting 9 points, grafting after curettage, and locking calcaneal plate placement as fixation material were planned.

Whether surgical treatment is required in intracalcaneal lipoma is still a controversial issue. However, these patients go for surgical treatment because they are radiologically like some other bone tumors and because of pain complaints. In surgery, the choice of grafting or cementing after curettage should be made in line with the surgeon's experience. The use of implants after grafting or cementation should be decided by evaluating both Mirels criteria and the criteria in the concept of critical size cyst.

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