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ABOUT COVER

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AIMS AND SCOPE

The primary aim of *World Journal of Clinical Cases* (WJCC, *World J Clin Cases*) is to provide scholars and readers from various fields of clinical medicine with a platform to publish high-quality clinical research articles and communicate their research findings online.

WJCC mainly publishes articles reporting research results and findings obtained in the field of clinical medicine and covering a wide range of topics, including case control studies, retrospective cohort studies, retrospective studies, clinical trials studies, observational studies, prospective studies, randomized controlled trials, randomized clinical trials, systematic reviews, meta-analysis, and case reports.

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Retrospective Study

Chronic otitis media and middle ear variants: Is there relation?

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Abstract

BACKGROUND

Chronic otitis media (COM) is an inflammatory disease that lasts for a long time. It is common in developing countries. Hearing loss can result from COM. The relationship between variations in middle ear anatomy and COM was investigated in our study.

AIM

To compare the prevalence of middle ear anatomic variations between the cases with COM and healthy individuals.

METHODS

This retrospective study included 500 patients with COM and 500 healthy controls. The presence of those variants was determined: Koerner's septum, facial canal dehiscence, high jugular bulb, jugular bulb dehiscence, jugular bulb diverticulum, sigmoid sinus anterior location and deep tympanic recesses.

RESULTS

A total of 1000 temporal bones were examined. The incidences of these variants were respectively (15.4%-18.6%), (38.6%-41.2%), (18.2%-4.6%), (2.6%-1.2%), (1.2%-0%), (8.6%-0%), (0%-0%). It was observed that only high jugular bulb ($P < 0.001$) and anteriorly located sigmoid sinus frequencies ($P = 0.002$) in the case group were statistically significantly higher than the control groups.

CONCLUSION

COM is a multifactorial disease and variants of middle ear have always been

important in terms of potential risk for complication during surgery but rarely associated with COM as an etiology or as a consequence of the disease. We didn't find a positive correlation between COM and Koerner's septum and facial canal defect. We ended up with a significant conclusion with the variants of dural venous sinuses -high jugular bulb, dehiscence of jugular bulb, diverticulum of jugular bulb and anteriorly located sigmoid sinus- that have been studied less and frequently associated with inner ear illnesses.

Key Words: Chronic otitis media; Radiology; Inflammation; Hearing loss; Mastoid cells; Head and neck

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Core Tip: Chronic otitis media (COM) is a chronic inflammatory disease. It is frequently seen in developing countries. COM can cause hearing loss. In our study, the relationship between the variations of middle ear anatomy and COM was investigated. Not many studies have been done on this subject in the literature. Our study was conducted by comparing patients and control groups. The relationship between different anatomical variations in the population and COM was investigated.

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INTRODUCTION

Chronic otitis media (COM) is a chronic inflammatory disease of the tympanic cavity and mastoid cavity mucosa that is characterized by a non-intact tympanic membrane and is a worldwide public health problem that primarily affects developing countries[1].

COM is classified clinically into two groups: Active and inactive, with multiple subgroups within each. This disease is one of the most common causes of conductive hearing loss; additionally, it can be progressive, with middle ear bones eroding and inner ear destruction leading to sensorineural hearing loss. Even though they are extremely rare, intracranial complications can occur[2].

Because of its high prevalence and the morbidities it causes, it plays an important role in the daily work routines of ear nose throat doctors and radiologists.

Chronic middle ear inflammation is caused by both genetic factors that cause functional and anatomic variations in the middle ear that predispose to COM and multiple environmental factors. Depending on the stage and complication of the disease, it is treated medically or surgically[3].

Middle ear anatomy is unique for every ear and person with multiple variations. High resolution computed tomography (HRCT) of the temporal bone provides plenty of information about the complications and the factors that predisposes to disease or to decide the kind of procedure to operate before surgery and to warn the surgeon against the risk of possible complications that might happen due to the variants with its high resolution and perfect bone detail.

The surgery procedure of choice for eliminating the infected mastoid cells is tympanomastoidectomy. Knowing the anatomic variations of that middle ear is essential before the operation in order to perform it properly; otherwise, recurrence and complications are unavoidable[4].

There hasn't been enough research done on the anatomic variations of the middle ear in the literature. The purpose of this study is to compare the prevalence of anatomic variations between case and control groups, as well as to define and compare the potential erosive effect of COM on the middle ear anatomic structures.

MATERIALS AND METHODS

This retrospective study included 500 patients with COM and 500 healthy controls. In both groups, the examinations were performed if a clinician deemed it necessary. Both groups were examined between March 2020 and August 2022.

Patients with maxillofacial anomalies, temporal bone fracture, prior ear surgery or mastoidectomy and younger than 18 years old were excluded from the study. Moreover, images which have low diagnostic quality due to motion artifacts were not included in the study.

We selected patients for our control group if COM was written as a final diagnosis in their electronic records and if there was mucosal thickening and soft tissue densities in middle ear space and mastoid cavity on imaging.

All images were examined by a radiologist who is 25 years experienced in head and neck radiology.

Seven variants were identified for review: Koerner's septum (KS), facial canal dehiscence, high jugular bulb, jugular bulb dehiscence, jugular bulb diverticulum, anteriorly located sigmoid sinus and deep tympanic recesses.

All studies were performed using a standardized temporal bone protocol with a revolution EVO scanner (General Electric Company, United States) with 100 kV (peak), 190 mA, 0.53:1 pitch, Display field of view (DFOV) 18.0 cm acquired at 0.625 mm section thickness.

To identify variants, the following criteria were used.

KS: KS was diagnosed on computerized tomography (CT) at the level of the antrum in the mastoid bone, with the appearance of a bone segment that is thicker and longer than the walls of air cells (Figure 1)[1].

Facial canal dehiscence: Facial nerve dehiscence is diagnosed when there is a loss of integrity in the bone channel (Figure 2)[5].

High riding jugular bulb: There are several criteria in the literature that indicate a high jugular bulb. We used the criterion that the roof of bulb reaches the cochlea's basal return level (Figure 3A and B)[6].

Jugular bulb dehiscence: When there is no lamina bone that separates the JB from the middle ear cavity and the bulb is in direct contact with the middle ear cavity, we diagnosed jugular bulb dehiscence (Figure 3C and D)[7].

Jugular bulb diverticulum: The jugular bulb diverticulum is a polypoid protrusion that extends superior to the bulb, as well as less medially and posteriorly (Figure 4)[7].

Anteriorly located sigmoid sinus: There must be no anteroposterior distance between the posterior part of the bony external auditory canal and the sigmoid sinus to diagnose the anteriorly located sigmoid sinus (Figure 5)[8].

Deep tympanic recess: If the depth of the recess is 6 mm or deeper on axial plane[9].

Statistical analyses

The Chi squared test and Fisher exact test (if necessary) were used to analyze categorical variables. The descriptive statistics were mean \pm SD, median [min max] for numerical variables and numbers and percentages (%) for the categorical variables. The SPSS Windows version 23.0 package program was used for the statistical analyses and *P* values less than 0.05 were considered statistically significant.

RESULTS

There are 500 patients in both control and case series. Case series has 260 male, 240 female patients and median age is 48.32 (\pm 15.03). Control series has 225 male, 275 female patients, median age is 45.86 (\pm 15.48)

In terms of clinical features, both groups were compared in Table 1.

The high jugular bulb frequency was found to be significantly different between the case (91 individuals; 18.2%) and control (23 individuals; 4.6%) groups (*P* < 0.001).

The incidence of anteriorly located sigmoid sinus was significantly higher in case groups (43 individuals; 8.6%) than in control subjects (19 individuals; 3.8%) (*P* = 0.002).

However, we didn't find a statistically significant difference in frequency with other variants.

In terms of the involvement sites of 500 cases diagnosed with COM and 500 healthy individuals are compared in Table 2.

There was a significant difference in facial canal dehiscence sides between case and control groups. (*P* = 0.003) While facial canal dehiscence is more common on the right side in case series (68, 35.4%), it has been observed that it is more common in the bilateral region in control series (*P* = 0.03).

DISCUSSION

Aside from clinic and audiologic tests, imaging modalities are used as a supplement to diagnose middle ear pathologies and plan surgery protocols. HRCT of the temporal bone is the modality of choice for its superior high resolution of bone anatomy.

It is critical for radiologists and otologists to understand temporal bone variants in order to understand the etiology and treatment of COM, which is a multifactorial disease.

Koerner's septum

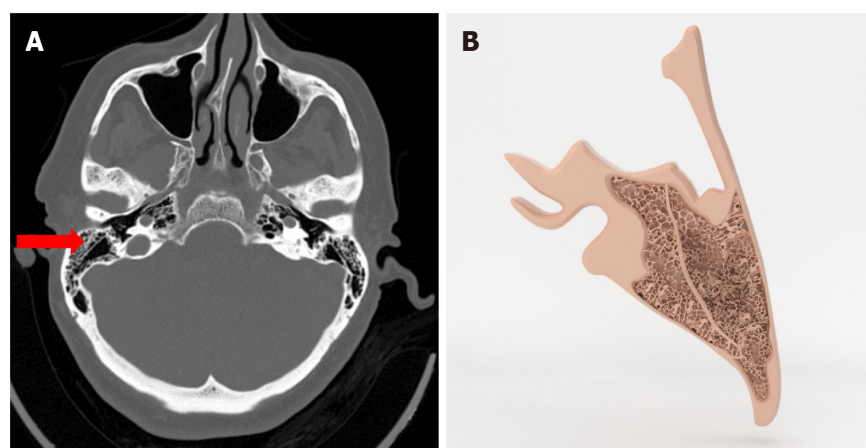
It is clinically significant that the air cells of the mastoid bone, which is formed by the union of squamous and petrous bones, are separated by a thin bone bridge named petrosquamous suture, as

Table 1 Comparison of the frequency of variants between case and control groups, n (%)

Variants ^a	Case (n = 500)	Control (n = 500)	P value
Koerner's septum +	77 (15.4)	93 (18.6)	0.178
Koerner's septum -	423 (84.6)	407 (81.4)	
Facial canal dehiscence +	193 (38.6)	206 (41.2)	0.401
Facial canal dehiscence -	307 (61.4)	294 (58.8)	
High jugular bulb +	91 (18.2)	23 (4.6)	< 0.001
High jugular bulb -	409 (81.8)	477 (95.4)	
Jugular dehiscence +	13 (2.6)	6 (1.2)	0.105
Jugular dehiscence -	487 (97.4)	494 (98.8)	
Jugular diverticulum +	4 (0.8)	0 (0)	-
Jugular diverticulum -	496 (99.2)	0 (0)	
Anterior located sigmoid sinus +	43 (8.6)	19 (3.8)	0.002
Anterior located sigmoid sinus -	457 (91.4)	481 (96.2)	
Deep tympanic recess +	0 (0)	0 (0)	-
Deep tympanic recess -	500 (100)	500 (100)	

^aP value was obtained from Pearson Chi Square test.

+: Indicates the presence of the variant; -: Indicates the absence of the variant.



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Figure 1 Koerner's septum in axial computed tomography image and illustration. A: Axial computed tomography image shows Koerner's septum (KS) as a thicker bone lamina in mastoid process (red arrow); B: KS is seen in illustration.

known as KS which is a common variant.

Previous researches in the literature have suggested that the KS inhibits pneumatization and drainage of the middle ear space by blocking aditus at the antrum and epitympanum. So it is thought as a reason of chronic middle ear diseases.

It's also a reason of complication in mastoid surgery[10,11].

Antrum, the largest air cell, is located deeper than the KS that's why it makes it more difficult to reach the antrum during surgery. Furthermore, during mastoid surgery, a well-developed, thick KS can be confused with the lateral wall of the sinus, and as a result, deep mastoid air cells cannot be cleaned out [10,11].

In our study, we found 77 patients (15.4%) with KS, and there are 93 cases (18.6%) with KS in the control series and statistically, it is not concluded to be significant with COM. Moreover we had more patients with this variant in our control group.

It is still important to report KS thus it is potentially a reason of complication in surgery.

Table 2 Comparison the frequency of the variants's side between case and control groups, *n* (%)

	Case (<i>n</i> = 500)	Control (<i>n</i> = 500)	<i>P</i> value
Koerner's septum			
Right	36 (46.8)	31 (33.3)	0.057
Left	29 (37.7)	34 (36.6)	
Bilateral	12 (15.6)	28 (30.1)	
Facial canal dehiscence			
Right	68 (35.4)	49 (23.8)	0.003
Left	84 (43.8)	85 (41.3)	
Bilateral	40 (20.8)	72 (35)	
High jugular bulb			
Right	39 (42.9)	10 (50)	0.763
Left	40 (44)	7 (35)	
Bilateral	12 (13.2)	3 (15)	
Jugular dehiscence			
Right	8 (61.5)	3 (50)	0.812
Left	4 (30.8)	2 (33.3)	
Bilateral	1 (7.7)	1 (16.7)	
Jugular diverticulum			
Right	3 (75)	0 (0)	---
Left	1 (25)	0 (0)	
Bilateral	0 (0)	0 (0)	
Anterior located sigmoid sinus			
Right	26 (60.5)	10 (62.5)	0.547
Left	14 (32.6)	6 (37.5)	
Bilateral	3 (7)	0 (0)	
Deep tympanic recess			
Right	0 (0)	0 (0)	---
Left	0 (0)	0 (0)	
Bilateral	0 (0)	0 (0)	

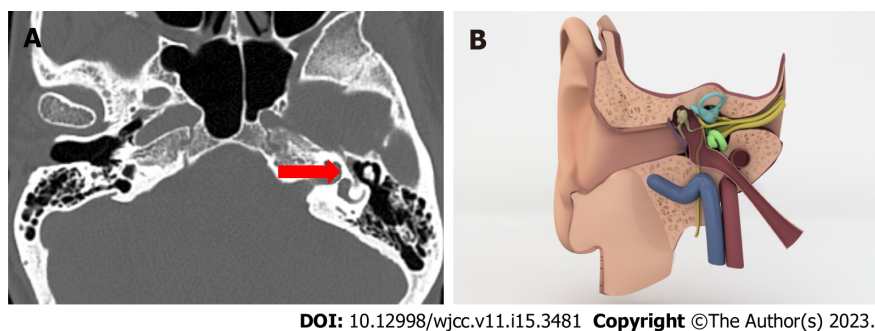
P value was obtained from Pearson Chi Square test.

Facial canal dehiscence

The chorda tympani runs through a bone labyrinth known as the fascial canal. It enters the temporal bone *via* the internal acoustic canal and exits through the stylomastoid foramen. It has three distinct parts along the course in the temporal bone. Respectively, labyrinth, tympanic and mastoid. Even though canal dehiscence can occur anywhere along the bonny canal, the tympanic segment is the longest and most common location[12].

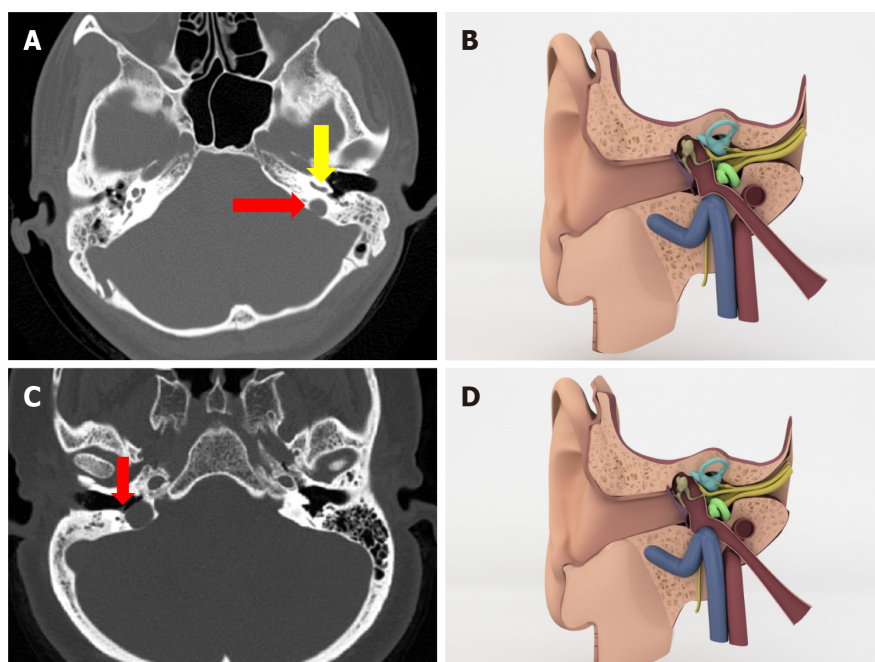
Facial canal dehiscence can be linked to ossification issues in inutero life. Despite the fact that it can be congenital, there are numerous studies that show a link between facial canal dehiscence and COM. It is probably a result of the erosive effect. That is why some consider it a pathology rather than a variant [13].

In imaging, a canal dehiscence should have precise edges so it can be differentiated from low mineralization. However micro dehiscence cannot be visualized on the temporal HRCT because of the thin structure and tortuous course of the canal. Which means, surgeons shouldn't depend on the CT report completely and should make a decent assessment of the canal intraoperatively in terms of preventing chorda tympani injury which may require revision operations[14].



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Figure 2 Labyrinth segment of facial canal without a bonny canal coverage in axial computed tomography image and illustration. A: Axial computed tomography image shows that there is not a bonny canal coverage around the labyrinth segment of facial nerve (red arrow); B: In this illustration labyrinth segment of facial canal can be seen without a bonny canal coverage.



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Figure 3 Axial computed tomography image and illustration. A: Axial computed tomography image shows that the roof of the jugular bulb (red arrow) is much higher than the basal turn of cochlea (yellow arrow); B: Illustration of middle ear cavity in coronal plane show jugular bulb colored as blue is slightly higher than the basal turn of cochlea; C: Axial computed tomography image shows that there is not a bone roof above jugular bulb and it is in direct contact with middle ear cavity (red arrow), tympanic membrane; D: When comparing with B we cannot see the bonny roof of jugular bulb.

In our study we have 193 patients in case series (38.5%) and 205 patients in control series (41%) with facial canal dehiscence. These numbers make it the most common variant in all of six variants we look for. We ended up with similar numbers so it is not statistically significant. So our results are more compatible with congenital theory.

Yet, it is essential to report facial canal dehiscence as inflammation of the middle ear can spread through the facial nerve and cause facial palsy[15] and its vulnerability in terms of getting harmed during surgery.

High riding jugular bulb

Jugular bulb is the venous part between the horizontal part of the sigmoid sinus and the upper end of the internal jugular vena. The morphology of the dural venous sinuses and the jugular bulb can vary greatly and is usually asymmetric for individuals[16].

Jugular bulb is absent at birth and develops around the age of two as a result of changing venous sinus dynamics after adapting to the erect position[17].

Although there is no agreement on the changing positions of sigmoid sinus and jugular bulb, general opinion is some pathologies that shorten the distance between the outer ear and the sigmoid sinus and rises the jugular bulb by reducing the pneumatization of the mastoid air cells[18].

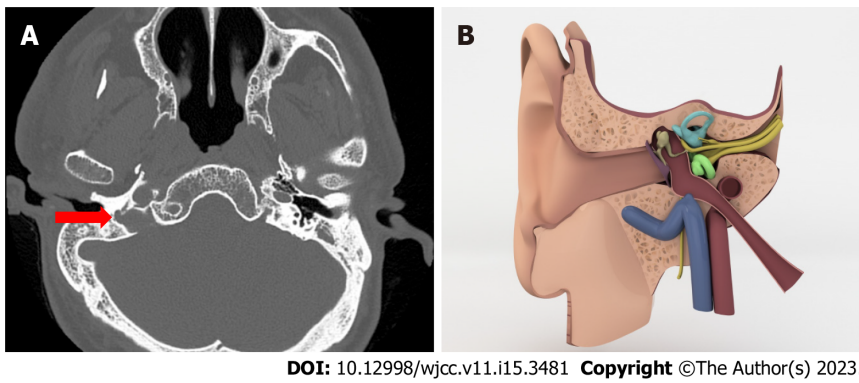


Figure 4 Jugular bulb diverticulum. A: Axial computed tomography image shows jugular bulb diverticulum as an outpouching from tiny bulbous (red arrow); B: Illustration draw of jugular bulb diverticulum which out pouches to middle ear cavity.

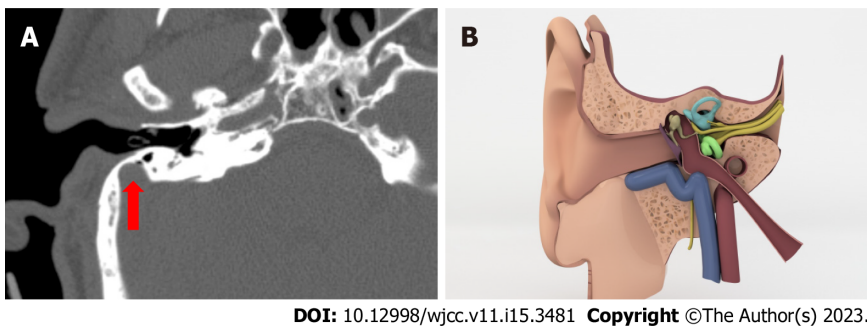


Figure 5 Anteriorly located sigmoid sinus. A: Anteriorly located sigmoid sinus. On axial CT images shows a sigmoid sinus very close to outer ear (red arrow); B: Illustration demonstrates anteriorly located sigmoid sinus.

The roof of the jugular bulb is lower anatomically than the floor of the hypotympanum, and a bone lamina should separate the jugular bulb from the middle ear space[18].

Authors suggested more than one criteria to decide if the jugular bulb is high. Any of them can be used. In this study we used the one which says if the jugular bulb is higher than the basal turn of the cochlea and projects into the middle ear space.

In the literature, the prevalence of the high riding jugular bulb, the most common variant of jugular bulb, is 3.5%-7.5%. It is mostly on the right side where the venous system is dominant and it is accepted as a physiologic variant.

In our study, 90 cases (18%) in the case series and 23 cases (4.6%) in the control series have a high riding jugular bulb. This time we found it statistically significant. We ended up with a prevalence in the control series that is comparable to the literature, but it is significantly higher in the case series. In the case series, 38 patients have a high riding jugular bulb on the right side, 40 patients have it on the left side and 12 patients have it bilateral. There is not a right side dominance unlike literature.

Jugular bulb dehiscence

Jugular bulb dehiscence is when jugular bulb doesn't have a bone lamina and is in contact with middle ear cavity. This condition creates a negative pressure effect in the cavity. This aggravates chronic middle ear inflammation and retracts tympanic membrane[19].

In our study, 13 patients (2.6%) have it, while 6 patients (1.2%) have it in the control series, indicating that there is no statistically significant relationship. However, it is worth noting that case series has twice as many people as case series.

Jugular bulb diverticulum

Jugular bulb diverticulum is considered as a true venous anomaly and is a reason of retrotympenic vascular mass. It can occur by two different entities. Either there is a dehiscence on the floor of middle ear wall or there is not a dehiscence but jugular fossa is large without a dehiscence floor[20].

We only have four patients in the case series and none in the control series, and there is no statistically significant relationship.

Anterior located sigmoid sinus

Anterior located sigmoid sinus is a rare variant like jugular diverticulum.

We have 43 patients (8.6%) in our case group and 17 patients (3.4%) in our control group. There are statistically significantly more patients in the case group than in the control group.

Jugular bulb and other venous variants are frequently associated with tinnitus, but they should not be considered as a cause of tinnitus before ruling out other possible etiologies because they could be often found in the general population. As far as we know, COM and jugular bulb and sigmoid sinus variants have never been associated in the literature before and our study is the first one.

Deep tympanic recess was not found in the total 1000 patients.

CONCLUSION

We aimed to find out a cause and effect relation between some of temporal bone variants and COM that's a multifactorial disease, characterized by the long term and destructive inflammation of the middle ear space and mastoid cellular. We didn't find a positive correlation between COM and KS and facial canal defect. There are more studies about both variants than others in the literature. On the other hand we ended up with a significant conclusion with the variants of dural venous sinuses -high jugular bulb, dehiscence of jugular bulb, diverticulum of jugular bulb and anteriorly located sigmoid sinus- that have been studied less and frequently associated with inner ear illnesses. Even though there is no agreement on why and how these variants occur, our study adds a new perspective to the literature by associating the variants with COM. However more studies should be done in the future to find out if these variants occur due to COM or COM occurs due to variants.

ARTICLE HIGHLIGHTS

Research background

Our article examines the anatomical and variational differences in chronic otitis media (COM) patients.

Research motivation

COM is a multifactorial disease and variants of middle ear have always been important in terms of potential risk for complication during surgery but rarely associated with COM as an etiology or as a consequence of the disease.

Research objectives

The objectives of our research are to determine its relationship with anatomical variations in COM.

Research methods

Our study is a retrospective study. The following conditions were investigated in our study: Koerner's septum (KS), facial canal dehiscence, high jugular bulb, jugular bulb dehiscence, jugular bulb diverticulum, anterior sigmoid sinus, and deep tympanic recesses.

Research results

Only the case group's high jugular bulb and anteriorly located Sigmoid Sinus frequencies were statistically significantly higher than the control groups.

Research conclusions

We discovered no link between COM and KS or facial canal defect. We came to a significant conclusion with the dural venous sinus variants -high jugular bulb, jugular bulb dehiscence, jugular bulb diverticulum, and anteriorly located sigmoid sinus- that have been studied less and frequently associated with inner ear illnesses.

Research perspectives

The relationship between COM and anatomical variations is investigated in our study.

FOOTNOTES

Author contributions: Gökharman FD and Şenbil DC conceived the idea for the study; Gökharman FD, Aydın S, Karavaş E, and Özdemir Ö designed and undertook the literature review; Yalçın AG, and Koşar PN collected data; Şenbil DC performed the statistical analysis, figures, and appendix and analyzed and interpreted the data; Gökharman FD and Aydın S wrote the first draft of the manuscript; Karavaş E and Özdemir Ö revised the subsequent drafts of the manuscript; All authors reviewed and agreed on the final draft of the manuscript.

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