



Association Between Mitral Valve Prolapse, Migraine, and White Matter Hyperintensities on Magnetic Resonance Imaging

Mitral Kapak Prolapsusu, Migren ve Manyetik Rezonans Görüntüleme Beyaz Cevher Hiperintensiteleri Arasındaki İlişki

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Abstract

Objective: Migraine is linked with an elevation in vascular risk factors, ischemic stroke, and a variety of constitutional brain lesions. However, the pathogenesis of this relationship is still inexplicit. The link between cardiac diseases and comorbid migraine-ischemic stroke might be a vascular disease involving both heart and brain. In this study, an association between mitral valve prolapse (MVP), migraine, and the presence of brain white matter hyperintensities (WMHs) were evaluated among adult subjects with migraine headache devoid of any traditional vascular risk factors.

Materials and Methods: Four hundred subjects (200 subjects with migraine headache, 200 healthy controls; age range 18-50 years) were incorporated in the retrospective study. Existence of a headache compatible with migraine was diagnosed according to the International Headache Society-2 criteria. The participants were devoid of any known comorbid diseases, vascular risk factors or inflammatory diseases. All patients, both those with migraine and controls were screened with echocardiography to assess for MVP and with brain magnetic resonance imaging to evaluate the presence of any WMHs.

Results: The prevalence of MVP was found to be higher in the migraine group ($p<0.011$). The odds ratio (OR) for the presence of MVP in patients with migraine compared with controls was 2.44 [95% confidence interval (CI): 1.25-4.74; $p=0.0086$]. The OR for the presence of WMHs in patients with migraine compared with controls was 5.88 (95% CI: 3.42-10.10; $p<0.0001$). After modifying for confounding factors, multiple linear regression analyses revealed that migraine was independently and positively associated with MVP ($p=0.044$), tricuspid regurgitation ($p=0.003$), and WMHs ($p<0.001$), and mitral regurgitation and migraine was independently and positively connected with WMHs ($p<0.005$ and $p<0.001$, respectively).

Conclusion: MVP is found to be independently associated with migraine when compared with controls. Therefore, we suggest that MVP might have an association with migraine. Nevertheless, we could not demonstrate any correlation between MVP and WMHs. Hence, we suggest that MVP might not be involved in the evolution of WMHs in migraine.

Keywords: Headache, vascular risk, cardiovascular risk, magnetic resonance imaging, echocardiography

Öz

Amaç: Migren, iskemik inme ve bazı yapısal beyin lezyonları gelişimi gibi vasküler risk faktörlerinde artış ile ilişkilendirilmektedir. Bununla birlikte bu ilişkinin patogenezi halen netleşmemiştir. Kardiyak hastalıklar ve iskemik inme ile komorbide gösterebilen migrenin bunlarla bağlantısı, hem kalp hem de beyin etkileyen vasküler bir hastalık olması şeklinde açıklanabilir. Bu çalışmada, geleneksel vasküler risk faktörleri olmayan genç-erişkin migren hastalarında mitral kapak prolapsusu (MVP) ile migren ve beyaz cevher hiperintensitesi (BCH) arasındaki ilişki incelendi.

Gereç ve Yöntem: Bu retrospektif çalışmaya 400 olgu alındı (200 migren ve 200 sağlıklı kontrol; yaş aralığı 18-50 yaş). Migren tanısı Uluslararası Baş Ağrısı Toplumu-2 kriterlerine göre kondu. Katılımcıların bilinen vasküler risk faktörleri, enflamatuvar veya eşlikçi başka hastalıkları yoktu. Migrenli kişiler ve sağlıklı kontrollere transtorasik ekokardiyografi yapılarak MVP varlığı; beyin manyetik rezonans görüntüleme yapılarak BCH olup olmadığı tarandı.

Bulgular: Migrenli kişilerde MVP prevalansı, kontrollere göre anlamlı yüksek ($p<0,011$) bulundu. Migrenli kişilerde MVP varlığı odds oranı (OR): 2,44 [%95 güven aralığı (GA): 1,25-4,74; $p=0,0086$] saptandı. Migrenli kişiler ve kontrol grubu BCH varlığı açısından karşılaştırıldığında OR oranı 5,88 (%95 GA: 3,42-10,10; $p<0,001$) bulundu. Karıştırıcı faktörlere göre uyarlandıktan sonra yapılan çoklu doğrusal regresyon analizleri migrenin, MVP ($p=0,044$), triküspid yetmezliği ($p=0,03$) ve BCH varlığı ($p<0,001$) ile bağımsız, pozitif korele olduğu; migrenin ve mitral yetmezliğin BCH açısından bağımsız, pozitif korelasyon gösterdiği gözlemlendi ($p<0,05$, $p<0,001$).

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Received/Geliş Tarihi: 29.01.2018 **Accepted/Kabul Tarihi:** 08.05.2018

Published in: The preliminary data of this work was presented as an oral poster at the 2015 World Neurology Congress.

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Turkish Journal of Neurology published by Galenos Publishing House.

Öz

Sonuç: MVP varlığı, kontrollerle karşılaştırıldığında migren ile bağımsız ilişkili bulundu. Bu nedenle MVP varlığının, migren ile ilişkili olabileceği düşünüldü. Bununla birlikte, MVP ile BCH arasında pozitif bir korelasyon bulunmadığı için MVP'nin, migrenli olgularda BCH gelişimine bir katkısı olmadığı düşünüldü.

Anahtar Kelimeler: Baş ağrısı, vasküler risk, kardiyovasküler risk, manyetik rezonans görüntüleme, ekokardiyografi

Introduction

Migraine is a very common neurovascular disorder caused by both hereditary (60%) and environmental factors (40%), affecting nearly 15% of the population (with a similar prevalence, 16.4% in Turkey) or about one billion people worldwide (1,2). It was reported to be the second leading cause of disability according to the annually performed "Global Years Lived With Disability" analysis in 2016 (3). Among its high prevalence and disability potency, migraine is also associated with an increase in vascular risk factors (both cardiovascular and cerebrovascular) according to risk prediction score for vascular diseases (4).

However, whether migraine itself is the causative factor or whether the disease is a marker of another underlying cause ending with cardio-and cerebrovascular diseases remains unclear (3,5,6). The suggested "higher prevalence of vascular diseases encompassing both heart and brain" might be due to an augmented presence of vascular endothelial dysfunction or due to a boosting inflammatory response to neurogenic inflammation, elevated prothrombotic factors or due to genetic predisposition. It might also be due to high consumption of anti-migraine medications. Also, the higher manifestation of vascular diseases among patients with migraine might be due to the "higher occurrence of constitutional heart anomalies" such as patent foramen ovale, atrial septal defects, pulmonary arteriovenous malformations, mitral valve prolapse (MVP), atrial septal aneurysm, and congenital heart disease (5,6,7).

MVP is the most frequently detected valvular condition affecting 2-3% of middle-aged individuals in the general population. Hence, MVP would be expected to affect 176 million people worldwide (8). According to some earlier studies, MVP was considered to be a disease with frequent and serious complications including stroke, atrial fibrillation, heart failure, and mitral regurgitation due to myxomatous degeneration, redundancy of the valve, and supraventricular arrhythmias (8,9,10). However, it was shown to be a benign entity, with a low occurrence of adverse events in a community-based FHS sample (11).

A variety of brain lesions, including white matter hyperintensities (WMHs), and clinically silent infarct-like lesions, localized especially in the posterior circulation have been associated with migraine. Hence, a meta-analysis indicated a four-fold increased risk for the occurrence of these WMHs in people with migraine compared with controls (12). Although these WMHs can also be detected in apparently healthy people, they are frequently regarded as ischemic in origin, and are associated with vascular risk factors. The determining factors in the development of these WMHs are considered to be the lifetime spent with migraine headaches and the frequency of attacks (13,14). However, it is also suggested that conditions as routine risk factors for stroke may lead to tissue damage via vascular endothelial dysfunction

or hypercoagulation or embolization, and these might then contribute to the evolution of these WMHs lesions (5,13,14).

There are many developments in the era of migraine pathophysiology and treatment: The underlying mechanism in migraine headache is proposed to be "triggering of the trigemino-vascular system," followed by liberation of calcitonin gene-related peptide" (CGRP) (15). Hence, many trials present strong verification that disturbance of the effects of endogenous CGRP is an effective commence to minimize migraine headaches. However, despite these developments in the pathophysiology of migraine, the mechanism by which it conveys an increased risk for the incidence of WMHs, even "silent infarct-like lesions," is still unclear (16).

We hypothesized that one of the factors responsible for the development of these WMHs in patients with migraine might be "the presence of MVP" in these patients (in case it exists as a comorbid condition). Therefore, we evaluated both the association of MVP with migraine, and the presence of WMHs in subjects with migraine with no traditional vascular risk factors.

Materials and Methods

Study Design and Patients

We analyzed data obtained from the records of patients admitted to our outpatient neurology clinic with a diagnosis of headache (compatible with migraine) from June 2014 till December 2016, retrospectively. The patients whose medical records contained both brain magnetic resonance imaging (MRI) and transthoracic echocardiography (ECHO) were included in the study.

Included patients with migraine headache were those with duration of symptoms of ≥ 1 year, repetition of headache ≥ 2 attacks/month, and absence of vascular risk factors along with absences of inflammatory or chronic illnesses such as renal, metabolic, infectious, or immune diseases. Patients with headache duration lasting < 1 year; those who used analgesics daily or migraine prophylaxis as propranolol, topiramate, or valproic acid; patients with a history of antecedent cerebrovascular or cardiovascular disease, arterial hypertension (blood pressure $> 130/80$ mmHg), diabetes mellitus, or hyperlipidemia; and those who were pregnant, breast feeding, or on hormonal contraceptives were excluded. Patients using drugs inclusive of antiplatelet agents, anticoagulants, statins, or hormonal drugs were also excluded. Patients who might have possible "symptomatic migraine headache" in whom MRI demonstrated ischemic infarcts, brain tumors, or other conditions that may be affiliated with migraine headache were excluded. Patients with any constitutional heart disease including patent foramen ovale, atrial septal defect, valvular heart disease other than MVP were also excluded. According to these inclusion and exclusion criteria, 200 consecutive patients diagnosed with migraine headache [143 women (71.5%), 57 men (28.5%); mean age 33.8 ± 7.6 (range: 18-

50) years] were recruited as the study group. For the control group, 200 healthy subjects [131 women (65.5%), 69 men (34.5%); mean age 33.7 ± 8.2 (range: 18-50) years] were convened from the hospital and laboratory staff, patients' relatives, and the general population. Inclusion criteria for the control subjects were absence of headaches such as migraine, tension-type or cluster headaches; absence of any neurologic or systemic diseases; and matching the patients with migraine in age. The exclusion criteria for the control subjects were the same as for the migraine study group.

Diagnoses of migraine headache were made as stated by the criteria of the International Classification of Headache Disorders-2 (17). Attack histories compatible with migraine were obtained in detail, subsuming disease duration (years), age at onset, repetition of attacks per month, average duration of headache (in hours), activating factors, accompanying manifestations including aura, site and seriousness of pain. The intensity of headache was appraised using a visual analogue scale (VAS) [range, 1 (minimum pain) to 10 (maximum pain)]. Patients were also requested to report the average VAS score. All patients and control subjects were examined thoroughly.

This study was evaluated by the Baskent University Institutional Review Board and due to its retrospective design it was exempted from the requirement for signed informed consent; however, oral informed consent was still taken from all patients.

MRI

Cerebral MRI (1.5 Tesla MR 355 Inspire; General Electric, SC, USA) with the orbito-meatal line as a reference were performed for all 400 participants. The scans comprised at least three sequences: sagittal T1-weighted, axial T2-weighted, and axial FLAIR images without intravenous contrast. The slice thickness was 5 mm with a gap of 1 mm. A radiologist who was blinded to the clinical details reviewed and scored all MRI scans.

The scans were evaluated for the existence and characteristics of WMHs, including apparition (focal or confluent), amount, measurements, dissemination (infra or supratentorial), and anatomic position. FLAIR images were used for the determination of the number and size of WMHs; focal punctate hyperintensities <10 mm were included, and large confluent hyperintensities (>9 mm) were excluded. No WMHs were detected in the corpus callosum. Periventricular WMHs were defined as those touching the ventricular system; juxtacortical WMHs were defined as those touching the cortical structures; all others were considered subcortical (18).

The subjects whom brain MRI revealed WMHs, were evaluated with laboratory tests for vasculitis (anticardiolipin antibodies, antinuclear antibody, lupus anticoagulant, anti-double-stranded DNA autoantibody, and C3 and C4 levels).

Echocardiographic Methods

All subjects underwent standard transthoracic ECHO (TTE) in accordance with the guidelines of the American Society of ECHO, which was performed by an experienced cardiologist who was blinded to any previous MVP diagnosis and clinical history (19). TTE was performed using a GE Health-Care Ultrasound Cardiology device containing a 3.5-MHz transducer. A diagnosis of MVP was made in case during systole, the view of parasternal long-axis displayed any superior displacement of mitral leaflets (one or more) exceeded 2 mm beyond a line attaching the annular

hinge points to the left atrium (20). At the same time, TTE was also used to rule out the existence of any valvular heart diseases other than MVP, atrial septal defects, patent foramen ovale, systolic dysfunction, or other constitutional heart diseases.

Statistical Analyses

We performed the statistical analyses with IBM SPSS for Windows version 22.0. The Shapiro-Wilk test was used for the evaluation of the normality of continuous variables. The Mann-Whitney U test was used for the assessment of differences between two groups, the independent samples t-test was used for continuous variables, and Pearson's chi-square or Fisher's exact test were used for categorical variables. Factors affecting migraine, MVP, and WMHs were identified using multiple logistic regression analyses. Significance was considered if a p value was less than 0.05. For the presentation of continuous variables, mean \pm standard deviation or median (range) were used. Categorical variables are reported as frequencies and percentages.

Results

The characteristics of the patients with migraine headache and characteristics of the control groups are depicted in Table 1. The odds ratio (OR) of MVP in patients with migraine headache was 2.44 [95% confidence interval (CI): 1.25-4.74; $p=0.086$], and the OR of WMHs in association with migraine was 5.88 (95% CI: 3.42-10.10; $p<0.001$). There was statistically significant difference between the sex of subjects with migraine because the number of female subjects was higher than for male subjects. Another statistically significant difference was found in the duration of migraine attacks with aura; the duration of attacks was longer than the duration of attacks without aura (Table 2). Except for these, there was no statistically significant differences between the characteristics of migraine headache, findings in ECHO or in the presence and localization of WMHs in migraine with and without aura group (Table 2).

After modifying for confounding factors, linear regression analyses disclosed that migraine headaches were independently and positively connected with MVP ($p=0.044$), tricuspid regurgitation ($p=0.003$), and WMH ($p<0.001$) (Table 3).

According to univariate analyses, MVP and mitral regurgitation were found to be significantly correlated ($r=0.482$, $p<0.001$). After regulating for confounding factors, multivariate logistic regression analyses revealed that mitral regurgitation was independently and positively correlated with MVP (Table 3). However, MVP was not significantly correlated with migraine characteristics or the presence or number of WMH (Table 3).

After altering for confounding factors, multivariate logistic regression analyses showed that WMH were independently and positively associated with female sex, age, migraine headaches, mitral regurgitation, and smoking compared with the normal brain MRI group (Table 3).

When the participants were subdivided into groups with and without WMH (Table 4), the presence of WMH was found to be significantly associated with older age, female sex, smoking, mitral regurgitation, tricuspid regurgitation, duration of migraine headache, and the frequency of migraine attacks (Table 4).

Table 1. Characteristics of the study participants with migraine and controls*

	Control (n=200)	Migraine (n=200)	P
Age (years)	33.7±8.2	33.8±7.6	NS
Sex (female/male)	131/69 (65.5%/34.5%)	143/57 (71.5%/28.5%)	NS
Ejection fraction	66.0±4.4	65.4±4.5	NS
Mitral valve prolapse	14 (7%)	31 (15.5%)	0.011
Mitral regurgitation	26 (13%)	44 (22%)	0.018
Tricuspid regurgitation	8 (4%)	31 (15.5%)	<0.001
Atrial regurgitation	-	8 (4%)	0.007
Presence of WMHs (%)	20 (10%)	79 (39.5%)	<0.001
Presence of juxtacortical WMHs	8 (40%)	48 (60.8%)	NS
Presence of subcortical WMHs	18 (90%)	77 (97.5%)	NS
Presence of periventricular WMHs	-	16 (20.3%)	-
Presence of infratentorial WMHs	-	14 (17.7%)	-
Number of WMHs	2 (1-6)	5 (1-15)	<0.001
Number of juxtacortical WMHs	0 (0-2)	1 (0-8)	0.017
Number of subcortical WMHs	2 (0-5)	3 (0-13)	0.010
Number of periventricular WMHs	-	0 (0-4)	0.029
Number of infratentorial WMHs	-	0 (0-3)	0.044
Aura	-	64 (32%)	-
Migraine disease duration (years)	-	5 (1-30)	-
Headache frequency (no/month)	-	3 (2-13)	-
Headache duration (h)	-	24 (1-96)	-
Visual analogue scale score	-	8 (4-10)	-
Smoking (≤pack/day)	33 (16.5%)	41 (20.5%)	NS

*Data are reported as number (%), mean ± standard deviation, or median (range).
WMHs: White matter hyperintensities, NS: Not significant (p>0.05)

Table 2. Characteristics of the migraine participants with and without aura*

	Migraine without aura (n=136)	Migraine with aura (n=64)	P
Age (years)	35.7±7.1	33.9±8.2	NS
Sex (female/male)	89/47 (65.5%/34.5%)	54/10 (84.3%/15.6%)	0.007
Ejection fraction	65.2±4.3	66±4.3	NS
Mitral valve prolapse	18 (13%)	13 (20%)	NS
Mitral regurgitation	29 (21.3%)	15 (23.4%)	NS
Tricuspid regurgitation	23 (16.9%)	8 (12.5%)	NS
Atrial regurgitation	6 (4.41%)	2 (3.1%)	NS
Presence of WMHs (%)	51 (37.5%)	29 (45.3%)	NS
Presence of juxtacortical WMHs	32 (62.7%)	16 (55.2%)	NS
Presence of subcortical WMHs	50 (98%)	27 (93.1%)	NS
Presence of periventricular WMHs	13 (25.5%)	3 (10.3%)	NS
Presence of infratentorial WMHs	7 (13.7%)	7 (24.1%)	NS
Number of WMHs	51 (1-6)	28 (1-5)	NS
Number of juxtacortical WMHs	31 (1-11)	16 (2-6)	NS
Number of subcortical WMHs	50 (1-13)	26 (1-7)	NS
Number of periventricular WMHs	13 (3-6)	3 (0-1)	NS
Number of infratentorial WMHs	7 (1-4)	7 (1-6)	NS
Migraine disease duration (years)	6 (1-30)	10 (1-30)	NS
Headache frequency (no/month)	5 (2-13)	4 (2-13)	NS
Headache duration (h)	19 (1-96)	30 (4-96)	0.001
Visual analogue scale score	7 (4-9)	8 (6-10)	NS
Smoking (≤pack/day)	27 (19.8%)	14 (21.8%)	NS

*Data are reported as number (%), mean ± standard deviation, or median (range).
WMHs: White matter hyperintensities, NS: Not significant (p>0.05)

Table 3. Factors affecting migraine, mitral valve prolapse, and white matter hyperintensities*

		Odds ratio	95% confidence interval	p
Migraine	Mitral valve prolapse	2.079	1.021-4.234	0.044
	Tricuspid regurgitation	3.583	1.542-8.327	0.003
	White matter hyperintensities	5.499	3.171-9.534	<0.001
White matter hyperintensities	Sex (female)	2.412	1.301-4.473	0.005
	Mitral regurgitation	2.395	1.306-4.391	0.005
	Migraine	5.740	3.265-10.090	<0.001
	Age	1.052	1.018-1.088	0.003
	Smoking	2.158	1.146-4.065	0.017

*Data are reported as the odds ratio and 95% confidence interval

Table 4. Characteristics according to the presence of white matter hyperintensities on magnetic resonance imaging*

	WMHs absent (n=301)	WMHs present (n=99)	p
Age (years)	33.1±7.9	35.6±7.7	0.006
Ejection fraction	65.7±4.5	65.8±4.3	NS
Sex (female/male)	107/194 (35.5%/64.5%)	19/80 (19.2%/80.8%)	0.004
Smoking (≤pack/day)	49 (16.3%)	25 (25.3%)	0.046
Mitral regurgitation	40 (13.3%)	30 (30.3%)	<0.001
Tricuspid regurgitation	23 (7.6%)	16 (16.2%)	0.022
Atrial regurgitation	4 (1.3%)	4 (4%)	NS
Mitral valve prolapse	30 (10%)	15 (15.2%)	NS
Aura	36 (29.8%)	28 (35.4%)	NS
Migraine disease duration (years)	4 (1-30)	10 (1-30)	0.024
Headache frequency (no/month)	3 (2-13)	4 (2-13)	0.011
Headache duration (hours)	24 (1-96)	24 (4-96)	NS
Visual analogue scale score	7 (5-10)	8 (4-10)	NS

*Data are reported as number (%), mean ± standard deviation, or median (range).
MRI: Magnetic resonance imaging, WMHs: White matter hyperintensities, NS: Not significant (p>0.05)

Discussion

A high prevalence of MVP and migraine comorbidity (as 20-59%) has been shown (21). Many brain MRI studies reported a relatively high prevalence of subclinical WMH in patients with migraine headache (13,14,22). Regarding these data, we wanted to evaluate the alliance relation MVP and migraine in conjunction with WMHs.

Our study also demonstrated that a higher number of patients with migraine headache encountered MVP and mild valve regurgitation compared with the control group (p=0.044). There was no statistically significant difference between MVP and migraine attacks with or without aura (Table 1, 2). The risk for MVP was 2.4-fold higher in patients with migraine than controls (OR: 2.44; 95% CI: 1.25-4.74; p=0.0086) (Table 3). Although mild mitral regurgitation was found to be independently associated with WMH, no association could be elicited between MVP and the presence of WMHs (Table 3).

Our results are in line with previous studies reporting MVP to be more frequent in subjects with migraine headache than in controls (21,23), although there are some conflicting results (24,25). The sample group was quite small in one study, and the mean age of the participants in another study was older than in our study, and no data were included on vascular risk factors (24,25).

In our study, the prevalence of MVP was found as 15.5% in subjects with migraine headaches, similar to the reported prevalence of MVP in the literature (15-25%) (8,23). A previous study reported that 27.8% patients with MVP had migraine headache, which was significantly higher than the expected number in the general population (approximately 10%) (26).

Although several mechanisms have been suggested regarding the causal association between the generation of migraine headaches and the development of MVP, the underlying mechanism is still obscure. It is suggested that shortened platelet survival and loss of endothelial continuity could initiate platelet activation and aggregation (21,23). As such, migraine headaches could be generated following the release of serotonin from these aggregated platelets (7,23). Paroxysmal arrhythmias that may accompany MVP syndrome may also increase the likelihood of embolization, leading to the formation of silent infarct-like lesions or WMHs (27).

Migraine headache is reported to be a risk factor for development of WMHs and the prevalence of WMHs in patients with migraine ranges from 14% to 59% (13,28,29). One suggested mechanisms for this is “oligemia triggered by repeated and prolonged periods of migraine attacks affecting the vulnerable small deep penetrating arteries” (13,28). In the present study, WMHs were detected 5.88 times more frequently in patients with migraine headaches than in controls, and we also showed that the presence of WMHs increased according to the (increased) headache frequency and duration, concordant with previous studies (Table 4). The association between WMHs and both the duration and frequency of migraine attacks suggest that the “migraine attack” itself plays a role in the pathophysiology.

Mitral regurgitation and other co-factors (such as age, female sex, mitral regurgitation, and smoking) are found to be independent factors associated with WMH (Table 3). Hence, these co-factors may also contribute to the pathophysiology of WMH.

Therefore, the relationship between mitral regurgitation and WMH still merits further investigation.

In this study, the anatomic location of WMHs differed between the groups. In subjects with migraine headache, WMHs were mostly located in the subcortical region, less juxtacortically; and only subjects with migraine headache had WMHs located in the periventricular and infratentorial regions (Table 1). These findings are in line with previous studies (13,28,29). However, in contrast to our findings, in another study no differences in the location of deep and periventricular WMHs could be elicited between migraineurs and controls (14). Accordingly, there may be different mechanisms affecting the development of these lesions in association with migraine. Hence, some previous studies included participants older in age with higher vascular risk factors (14). There are also studies reporting no increased number of WMHs in association with migraine compared with controls (30,31), but those studies had either smaller sample numbers than ours or included no information about the vascular risk factors of their participants (30,31).

MVP is a common decadence cardiac valvulopathy of unknown etiology inclining to mitral regurgitation, heart failure, and even sudden fatality. Subjects with MVP have a greater chance to experience mitral regurgitation than those without, even if the valvular regurgitation is trace or mild (11,32). The prevalence of mild mitral regurgitation was 68.9% in our study, consistent with previous reports (11). The most common complication of MVP is worsening of the mitral regurgitation. It causes diastolic dysfunction of the left atrium and left ventricle, cardiac arrhythmias, and subsequent heart failure, with an increased risk for infective endocarditis. Hence, 2.4% of patients with MVP experience thromboembolic events (33).

As far as we know, our study is the first to display a connection between MVP and WMHs in migraine. The unique feature of our study design was the simultaneous assessment of an association between MVP and WMH in migraine. In order to minimize the effects of age-dependent WMHs, we included patients aged <50 years who were devoid of any vascular risk factors or inflammatory diseases.

Study Limitations

There are many limitations of our study, such as its retrospective design and lack of trans-esophageal ECHO examination, because it is a better tool for diagnosing MVP. Furthermore, we could not strictly exclude risk factors such as smoking and obesity; we excluded heavy smokers (>1 pack/day), but did not exclude all cigarette smokers. However, there were no significant differences in smoking habits between the subjects with migraine and controls. We did not calculate the body mass indexes of the participants, which might also give more information. The association between migraine, WMHs, and MVP might have been evaluated further if we could have studied markers of thrombosis and coagulation factors in detail.

Conclusion

We found that MVP was independently associated with migraine headache, but we found no correlation between MVP and WMHs. We suggest that MVP is not causally involved in

the development of WMHs associated with migraine headache. Further prospective studies should evaluate the relationships between migraine, WMHs, and MVP.

Acknowledgements

The authors thank Hakan Gökahmetoğlu for helping with data collection.

Ethics

Ethics Committee Approval: This was a retrospective study, no ethical committee approval was taken. All procedures performed in studies involving human participants were in accordance with the ethical standards of the institutional and/or national research committee and with the 1964 Helsinki Declaration and its later amendments or comparable ethical standards.

Informed Consent: This study was deemed to be exempt from the requirement for informed consent by the Institutional Review Board of Baskent University due to its retrospective design.

Peer-review: Externally peer-reviewed.

Authorship Contributions

Concept: A.Y.A., M.K.T., Design: A.Y.A., M.K.T., Data Collection or Processing: A.Y.A., M.K.T., H.L., S.A., Analysis or Interpretation: A.Y.A., Literature Search: A.Y.A., M.K.T., Writing: A.Y.A., M.K.T.

Conflict of Interest: The authors declare that they have no conflict of interest.

Financial Disclosure: The authors declared that this study received no financial support.

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