Is Idiopathic Benign Paroxysmal Positional Vertigo Associated with Serum 25-Hydroxy Vitamin D Deficiency?

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Objective: Benign paroxysmal positional vertigo (BPPV) is a common cause of peripheral vertigo in the general population. We investigated the role of 25-hydroxy (25-OH) vitamin D deficiency in BPPV by comparing 25-OH vitamin D levels in healthy controls and in patients with BPPV.

Methods: 25-OH vitamin D levels of 125 patients with idiopathic BPPV who were diagnosed at our clinic between January 2018 and September 2018 and 101 healthy controls without vertigo were compared statistically.

Results: In this study, vitamin D deficiency was detected in patients diagnosed with BPPV, but there was no statistically significant difference with the control group.

Conclusion: The prevalence of the vitamin D deficiency is very high in our population. Despite the major studies in the literature, vitamin D deficiency was not related to BPPV as a result of this research.

Keywords: Benign paroxysmal positional vertigo, 25-hydroxy vitamin D, peripheral vertigo

INTRODUCTION

Benign paroxysmal positional vertigo (BPPV) is the most common cause of peripheral vertigo in the general population (1). BPPV is characterized by vertigo that is triggered by head movements and lasts for seconds, and accompanied by a feeling of imbalance and nausea. In the pathophysiology of BPPV, there are two theories called cupulolithiasis and canalithiasis. There are three semicircular canals located perpendicular to each other in the inner ear and sensing the angular movements of the head: posterior, lateral (horizontal), superior (anterior) canal crura are associated with the utricle, known as autolytic organ. Calcium (Ca) carbonate crystals are found in the otocional layer above the maculae found in the utricle. The otocinia separated from the utricle macula can pass into semicircular canals. Vertigo and nystagmus can occur when these otocinia stimulate cupula. The “canalithiasis” theory suggests that the free movement of these otocinia in the canal plays a role in the pathophysiology of the disease. It was first described by Hall, Ruby and McClure in 1979 and was first proven in vivo by Parnes and McClure in 1992. “Cupulolithiasis”, defined by Schuknecht in 1969, refers to the adherence of the otoconia to the cupula (2). Otoconia contains Ca carbonate in the form of Ca crystals and an organic core consisting mainly of glycoproteins. Ca metabolism also plays a primary role in the synthesis/absorption of otocinia and is therefore theoretically thought to be an etiological factor at the onset of BPPV (3). The aim of our study was to compare 25-hydroxy (25-OH) vitamin D levels in patients with idiopathic BPPV and healthy controls, and to investigate the role of 25-OH vitamin D in the development of BPPV.

METHODS

One hundred and twenty-five (100 female and 25 male, mean age=52±14 years) patients, who were admitted to the vertigo outpatient clinic between June 2018 and September 2018 and were diagnosed with idiopathic BPPV, were included in this retrospective case-control study. The patients had no history of Meniere’s disease, vestibular migraine, labyrinth hypofunction, head trauma or other vestibular diseases. The control group
consisted of 101 (74 female and 27 male, mean age=48±13 years) healthy volunteers. The patients had no history of neurological symptoms or vestibular disease. Patients with neurotologic symptoms and complaints of dizziness and imbalance were excluded from the study. All participants did not receive Ca or vitamin D treatment within the last year.

Vestibular evaluation was performed using computerized (videonystagmography: ICS Charter EP, GN Otometrics, USA). BPPV was diagnosed by Dix-Hallpike and Pagnini-McClure maneuvers. There was posterior canal involvement in 80 patients (64%), horizontal canal involvement in 38 patients (30.4%) and anterior canal involvement in seven patients (5.6%). Epley maneuver was used in patients with posterior canal involvement and Barbecue maneuver was applied to those with horizontal canal involvement. In the anterior canal involvement, “reverse Epley maneuver” was performed. Canalithiasis was detected in 72% (90 patients) of these patients and cupulolithiasis was responsible for the pathophysiology in 28% (35 patients) (Table 1). Blood was collected from patients with BBPV and healthy volunteers and 25-OH vitamin D levels were measured. 25-OH vitamin D levels were classified as normal (≥30 ng/mL), insufficient (>20 to <30 ng/mL) and deficiency (≤20 ng/mL). The approval of the Ethics Committee was obtained (dated: 6.11.2018, numbered: 48670771-514.10). Informed consent was obtained from the patients.

### Statistical Analysis

Statistical analysis was performed using SPSS version 23.0. Descriptive data are presented using mean and standard deviation for normally distributed variables, and median, minimum and maximum values for non-normally distributed variables (and frequency tables for ordinal variables). Chi-square was used to compare categorical variables. The suitability of the measured variables to normal distribution was examined by visual (histogram) and analytical methods (Kolmogorov-Smirnov/ Shapiro-Wilk tests). Pairwise comparisons were performed using Student’s t-test for normally distributed parameters and Mann-Whitney U and Kruskal-Wallis tests for non-normally distributed parameters. P<0.05 was evaluated as statistically significant.

### Results

The mean serum 25-OH vitamin D levels were 16.36 ng/mL (3.52-53.91) in the BBPV group and 17.09 ng/mL (4.46-53.51) in the control group. Vitamin D levels were low in both groups. In the BBPV group, 81 patients (65.3%) had serum 25-OH vitamin D deficiency, 33 patients (26.6%) had insufficient and 10 patients (8.1%) had normal levels. In the control group, 74 patients (73.3%) had serum 25-OH vitamin D deficiency, 16 patients (15.8%) had insufficient and 11 patients (10.9%) had normal levels (Table 2). There was no statistically significant difference between the BBPV

### Table 1. Rate of semicircular canal involvement and 25-hydroxy vitamin D levels

| Affected canal | n | % | Median (minimum-maximum) | Vitamin D levels | H | p* |
|---------------|---|---|--------------------------|-----------------|---|----|
| Superior      | 7 | 5.6 | 14.52 (9.60-36.04) | 0.645 | 0.724 |
| Posterior     | 38 | 30.4 | 18.00 (3.52-53.91) |                  |   |    |
| Lateral       | 80 | 64 | 16.18 (6.12-49.78) |                  |   |    |

*Kruskal Wallis-H test

### Table 2. Comparison of 25-hydroxy vitamin D levels between benign paroxysmal positional vertigo and control group

| n | BPPV (n=125) | Control group (n=101) | Significance |
|---|--------------|-----------------------|--------------|
| Gender | % | n | % | x² | p |
| Male  | 25 | 20.0 | 74 | 26.7 | 1.429 | 0.232 |
| Female | 100 | 80.0 | 74 | 73.3 | 0.526 | 0.468 |
| Age* | 52±14 | 48±13 | t=1.903 | 0.058 |
| Vitamin D groups | % | n | % | x² | p |
| Deficient (≤20 ng/mL) | 81 | 65.3 | 74 | 73.3 | 3.952 | 0.139 |
| Insufficient (>20 to <30 ng/mL) | 33 | 26.6 | 16 | 15.8 | 0.526 | 0.468 |
| Normal (≥30 ng/mL) | 10 | 8.1 | 11 | 10.9 | 0.526 | 0.468 |
| Vitamin D groups | % | n | % | x² | p |
| Deficient (<30 ng/mL) | 114 | 91.9 | 90 | 89.1 | 0.526 | 0.468 |
| Normal (≥30 ng/mL) | 10 | 8.1 | 11 | 10.9 | 0.526 | 0.468 |

BPPV: Benign paroxysmal positional vertigo, *Mean ± standard deviation values are given, **Since vitamin D values do not show normal distribution, median (minimum-maximum) values are given, Pearson chi-square test, Independent samples t-test, Mann-Whitney U test
group and control group in terms of 25-OH vitamin D deficiency (p=0.139). There was no difference between 25-OH vitamin D levels in BPPV patients regarding affected canals (p=0.724).

DISCUSSION

BPPV is the most common cause of peripheral vertigo at any age. The mechanism of BPPV is explained by the passage of the otoconia separated from the utricle into semicircular canals. There is no consensus on the factors that cause BPPV. Since the etiologic factors are unclear, most cases are considered idiopathic. Predisposing factors are senility, female gender, hormonal factors and viral causes (4). Inner ear consists of cochlea and labyrinth system. The bony labyrinth consists of three semicircular canals: superior (anterior), posterior and horizontal (lateral). There is a fluid called perilymph in the bony labyrinth. Membranous labyrinth consists of utricle, sacculle and membranous semicircular canals, and it contains endolymph. Membranous semicircular canals are located perpendicular to each other. The dilated parts are called ampulla. There are special cells in this region called “crista ampullaris” that is the sensory organ of balance. Within the wall of the utricle, there are cells called the macula of utricle, which lie horizontally and receive the sense of balance, and supporting cells. These cells have Ca$^{2+}$ particles called otoconia. Otoconia crystals have central and peripheral portions. The core is predominantly organic with a lower Ca$^{2+}$ level and the periphery is largely inorganic with a higher Ca$^{2+}$ level (5). Endolymphatic Ca$^{2+}$ concentration is critical for normal auditory and balance system (6-8). According to theoretical considerations, a link between otolytic disorders and vitamin D deficiency is highly probable. Endolymphatic (cochlea 23 μM and vestibule 280 μM) Ca concentration is much lower than perilymph. Yamauchi et al. (9) first demonstrated the expression of a complete Ca$^{2+}$ absorptive system in cochlear and peripheral portions. The core is predominantly organic with a lower Ca$^{2+}$ level and the periphery is largely inorganic with a higher Ca$^{2+}$ level (5). 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The relationship between idiopathic BPPV and vitamin D deficiency is controversial in the literature. As in our country, vitamin D deficiency is common in populations with short and variable sun exposure. We found low levels of vitamin D in both study and control groups. In this respect, further studies are needed to investigate the relationship between BPPV and vitamin D deficiency.

CONCLUSION

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