Update on benign paroxysmal positional vertigo

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Received: 15 October 2020 / Revised: 4 November 2020 / Accepted: 8 November 2020 / Published online: 24 November 2020

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Abstract
Benign paroxysmal positional vertigo (BPPV) is the most common cause of vertigo worldwide. This review considers recent advances in the diagnosis and management of BPPV including the use of web-based technology and artificial intelligence as well as the evidence supporting the use of vitamin D supplements for patients with BPPV and subnormal serum vitamin D.

Keywords
Dizziness · Vertigo · Benign paroxysmal positional vertigo

Introduction
Benign paroxysmal positional vertigo (BPPV) is characterized by paroxysms of vertigo triggered by head position changes in the direction of gravity [1]. BPPV is explained by migration of degenerated otoconia into the semicircular canals, rendering them sensitive to head motion [2]. BPPV is the most common cause of dizziness/vertigo worldwide with a lifetime prevalence of 2.4%, a 1-year prevalence of 1.6%, and 1-year incidence of 0.6% [3]. BPPV accounts for 24.1% of all hospital visits due to dizziness/vertigo [4]. BPPV is most common in elderly women with a peak incidence in their sixties and a women-to-men ratio of 2.4:1 [4]. Recurrences of BPPV are frequent [5, 6] with an annual recurrence rate of 15–20% [7, 8].

Even though benign in nature, patients with BPPV are markedly limited in their daily activities [10, 11]. The medical costs for diagnosis of BPPV have been estimated at 2,000 US$ in the USA [12], 364 Euros (~450 US$) in Spain [13], RMB 4165.2 Yuan (~600 US$) in China [14], and 180 US$ in South Korea [9]. Therefore, the healthcare burden due to BPPV totals around 2 billion US$ in the USA per year [15] and is likely to increase as the population ages. According to data from South Korea, the number of hospital visits per 100,000 of general population due to dizziness and vertigo was around 3974 in 2019 and could increase to 6057 by 2050, which corresponds to an increase of 52% [4].

Pathophysiology
The cause of BPPV is mostly unknown although cases may be associated with head trauma, a prolonged recumbent position, or various disorders involving the inner ear [16]. Other risk factors for BPPV may include female gender, age under 65 years, a high income, living in a metropolis, osteoporosis, hypertension, and non-apnea sleep disorders [17–20]. Indeed, a recent meta-analysis found that female gender, hypertension, diabetes mellitus, hyperlipidemia, osteoporosis, and vitamin D deficiency are risk factors for recurrences of BPPV [19]. Linked to this is the finding that elderly women with a lack of physical activity have a 2.6 times higher risk for BPPV than those who undertake regular activity [21]. Furthermore, vitamin D deficiency (<10 ng/ml) and insufficiency (10–20 ng/ml) have been associated with BPPV with an odds ratios of 3.8 and 23.0 [22]. This has obvious therapeutic implications (see below).
Clinical features

In the western countries, the posterior canal (PC) has been known to be the most commonly involved (88–90%) in BPPV with a predilection for the right ear [23, 24]. In studies conducted in South Korea, however, involvement of other semicircular canals (not just the PC) has been shown to be more common than previously reported in western countries [25]. Even though the PC was most commonly involved (59–61%), BPPV of the non-pure PC-BPPV type comprises about 40% of total BPPV, especially with involvement of the horizontal canal (HC-BPPV), where the otolithic debris may be located either in the canal (canalolithiatic or geotropic, 61–66%) or attached to the cupula (cupulolithiatic or apogeotropic, 31–33%) [10, 21]. The causes for this discrepancy is unknown [25].

Diagnosis

The International Classification of Vestibular Disorders (ICVD) formulated by the Barany Society established the diagnostic criteria for BPPV [26], which include recurrent attacks of positional vertigo/dizziness provoked by position changes, and the characteristic positional nystagmus elicited by each positional maneuver according to the subtype and affected ear [26]. During the Dix–Hallpike maneuver for diagnosis of PC-BPPV, a pillow may be placed under the shoulders instead of extending the patient’s neck about 30° below the table [27]. This modified maneuver may be useful in the limited clinical setting or in patients with a limited range of motion or difficulty relaxing their neck. HC-BPPV is diagnosed by determining the direction and relative intensity of the horizontal nystagmus induced during head-turning while supine. In both the canalolithiatic (geotropic) and cupulolithiatic (apogeotropic) types of HC-BPPV, the nystagmus beating to the lesion side is greater than that to the healthy side [1]. The accuracy of bedside lateralization of the affected side is acceptable in HC-BPPV when the nystagmus asymmetry is more than 30% [28]. When the intensities of nystagmus triggered during supine head-rolling test are similar between the directions, the direction of nystagmus induced by lying down or head bending (bow and lean test) may aid in lateralization of the involved side [29, 30]. The lying-down nystagmus mostly beats away from the affected ear in geotropic HC-BPPV but beats toward the affected ear in the apogeotropic type, while the reverse holds for the head-bending nystagmus [29, 30]. Patients with PC-BPPV may show vertical nystagmus during the bow-and-lean test [31]. A recent study found that head-shaking nystagmus (the nystagmus observed after head oscillation at 2–3 Hz in the horizontal plane for 20 cycles) occurs in about a half of the patients with HC-BPPV in the direction of head-bending nystagmus, thus helping lateralize the affected side [32].

The duration of vertigo and nystagmus is typically less than one minute in the canalolithiatic type of HC-BPPV [33]. Nevertheless, patients occasionally show persistent geotropic nystagmus while supine head-turning. This persistent geotropic positional nystagmus may be observed in association with focal central lesions [34], alterations in the specific gravity of the cupula or endolymph (light cupula) [35–37], or in migraine [38, 39] even though the mechanisms remain to be elucidated. Paroxysmal downbeat nystagmus may be observed during head extension not only in central lesions or BPPV involving the anterior canal (AC-BPPV) but due to compression of both vertebral arteries [40]. Positional nystagmus due to central lesions (central positional nystagmus, CPN) may be either paroxysmal or persistent, and both types of CPN may be ascribed to impaired central processing of canal and otolith signals [41]. Even though CPN may be differentiated from BPPV based on the temporal patterns of nystagmus intensity, occurrence of nystagmus in multiple planes, and additional ocular motor or other neurological findings indicative of central lesions [41, 42], clinicians should be careful when the seemingly BPPV does not respond to repeated canalith repositioning procedures.

Otolin-1 is an inner ear-specific collagen that forms a scaffold to promote optimal formation of the otoconia. Owing to its potential passage through the labyrinth-blood barrier, otolin-1 can be detected in the peripheral blood and may serve a biomarker for BPPV [19]. Thus, high serum levels of otolin-1 (> 300 pg/ml) may discriminate patients with BPPV from healthy controls [20].

The gold standard for diagnosing and determining the subtype of BPPV is the observation of characteristic nystagmus triggered during the positional maneuvers as mentioned above. However, a few studies have explored the utility of questionnaires in confirming BPPV and determining the subtypes [43–45] based on the characteristics (positional triggering, duration etc.) of the vertigo and positional changes that mostly induce it. [43]. A recent study investigating this questionnaire approach showed an acceptable sensitivity and specificity for the diagnosis of BPPV [45]. The questionnaire is comprised of six questions. The first three are designed to diagnose BPPV, and the latter three to determine the subtype and affected ear. This new approach for self-diagnosis of BPPV using a questionnaire would pave the way for self-administration of an appropriate canalith repositioning procedure when BPPV occurs and recurs.

By virtue of both recent developments in information (IT) and biology (BT) technology through programs available
on mobile devices [46] and using artificial intelligence and a deep-learning model, interest has turned to whether this approach can be used to determine the underlying disorder(s) causing dizziness and vertigo, and with this the subtype of BPPV [47]. Furthermore, recording of nystagmus during the attacks of vertigo may also become feasible in near future using various portable devices [48]. These kinds of approaches adopting artificial intelligence, deep-learning, wearable devices, and mobile applications, will become more important in determining the cause of vertigo, especially when we have to rely more on telemedicine as is the case now given the COVID-19 pandemic.

BPPV patients show 1-year recurrence rates of approximately 20% and 5-year recurrence rates of approximately 50% [44]. Repeat attacks of BPPV are frequent [5, 6] with the recurrences in a half of the patients within 40 months [7, 8]. A recent study found that these recurrences were somewhat random with only 24% of them involving the same canal on the same side as was affected in the previous attack [9]. A role for vitamin D in BPPV has recently been shown to be significant. A recent randomized clinical trial, patients with BPPV were randomly allocated either to have measurements of serum vitamin D and take 400 IU vitamin D and 500 mg calcium twice a day when the serum vitamin D level was subnormal or to be simply followed-up for the recurrences without intervention. A significant reduction in the recurrences was found in the treatment group (Fig. 1a) [73]. A subsequent meta-analysis of six randomized trials also proved the preventive effect of vitamin D supplementation on recurrences of BPPV (Fig. 1b) [74]. Thus, supplementation of vitamin D should be considered in patients with recurrent BPPV and subnormal serum vitamin D.

Otolith dysfunction could be associated with an increase in the recurrences of BPPV [75, 76]. Thus, the vestibular rehabilitation program including may reduce the recurrence rate of BPPV [77]. This program includes jumping on the trampoline-like surface with eyes open and closed, reading a text during linear head movements, standing on the tilt board and using an exercise ball.

**Conclusion**

Even though benign, BPPV brings with it major socio-economic impacts. Being able to better diagnose this, especially using remote devices would be a major breakthrough and ways to do this are being actively explored with encouraging data to support their adoption. In addition, while a range of physical therapies can be successfully used to treat BPPV,
there are interesting new data suggesting a role for vitamin D supplementation in a subgroup of patients with low serum levels of this vitamin.

Acknowledgements Funding was provided by Korea Healthcare Technology R&D Project, Ministry of Health and Welfare (Grant no. HI16C0429).

Compliance with ethical standards

Conflicts of interest Dr. H-J Kim, and Dr. J-H Park report no disclosures. Dr. J-S Kim serves as an Associate Editor of Frontiers in Neuro-otology and on the editorial boards of the Journal of Clinical Neurology, Journal of Neuro-ophthalmology, Journal of Vestibular Research.

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