Introduction

Benign paroxysmal positional vertigo (BPPV) is one of the most common diseases that can cause dizziness. Its prevalence is reported to be 11–64 per 100000 people, with a reported lifetime prevalence of 2.4%.

Although BPPV can be resolved with appropriate canalith repositioning procedures (CRPs), it also causes great mental stress in patients and increases the risk of falling, further jeopardizing the health of the afflicted individual.

BPPV occurs when otoconia are detached from the otolithic organ and enter the semicircular canal (SCC). Although it is suspected that otolith organ degeneration is the cause of otoconia detachment, the pathophysiology of this detachment is not clearly identified in most BPPV patients. However, in some cases, the cause of otoconia detachment from the otolithic organ is readily apparent. For example, in case of BPPV...
occurring after head trauma, especially in patients with temporal bone fracture, the cause is relatively clear. Because the otolith organ is located within the temporal bone, it is subjected to a large external impact in the event of temporal bone fracture. As a result, this may destroy the filaments attaching the otoconia to the otolith organ. Consequently, the otoconia may be detached from the otolith organ resulting in BPPV. Moreover, we believe that BPPV caused by temporal bone fracture would have greater damage to otolith organs relative to idiopathic BPPV. Accordingly, it was estimated that more otoconia would detach from the otolith organs, and that reattachment would be difficult after the CRPs. Therefore, we hypothesized that BPPV accompanied by temporal bone fracture would be more resistant to CRPs than idiopathic BPPV.

The incidence of BPPV in head trauma has been reported to be between 4.1% and 14.9%. Therefore, the incidence of BPPV is also expected to be high after temporal bone fractures. Despite the expected high incidence, few studies have been done on BPPV accompanied by temporal bone fractures. In this study, we aimed to investigate whether temporal bone fracture affects the number of CRPs performed for the remission of BPPV using univariable and multivariable regression analysis. The characteristics of BPPV accompanied by temporal bone fractures can be identified and, ultimately, these data could contribute to the identification of BPPV pathophysiology.

Subjects and Methods

Study design, population and study setting

This study was conducted with the approval of Institutional Review Board of our hospital (IRB No. 1912-013-16292). This retrospective study included patients diagnosed with BPPV of the canalolithiasis type between January 2016 to December 2018, at our tertiary hospital. BPPV was diagnosed based on the presentation of typical whirling-type dizziness on positional changes and typical nystagmus during specific maneuvers, such as the Dix-Hallpike, and head-roll test identified by videonystagmography. Appropriate CRPs, such as Epley or barbecue rotation maneuvers, were conducted according to the affected SCC. In a subsequent examination, the patient’s symptoms and nystagmus were assessed, and if BPPV persisted, the appropriate CRP was performed. Otherwise, the CRP was not performed. The number of CRPs performed until the remission of BPPV was counted. To consistently estimate the number of CRPs, patients with an interval between CRPs for BPPV other than 3 or 4 days were excluded.

Fig. 1. The representative CT images of temporal bone fractures.
The presence of temporal bone fracture was confirmed by temporal bone CT (Fig. 1). Among the patients with temporal bone fracture, patients with otic capsule-disrupting fracture were excluded. Among the patients without temporal bone fracture, patients with factors that could induce BPPV such as head trauma were excluded. Patients with multiple canal involvement or canal conversion during CRPs were excluded. Clinical features of the patients and information about BPPV were collected from their medical records.

Statistical analysis

Age, male and female sex ratios, number of CRPs performed for the remission of BPPV, prevalence of disease, and the side and type of the affected SCC were compared between patients with and without temporal bone fracture using the Mann-Whitney, Pearson chi-square or Fisher’s exact test. To identify potential factors related to the number of CRPs, individual factors, including sex, age, the presence of temporal bone fracture, hypertension, diabetes, dyslipidemia or osteopenia, BPPV side, and the type of affected SCC were analyzed using univariable regression analysis. Then multivariable regression analysis was performed using only the variables with a p-value of 0.2 or less in the univariable analysis. Data were analyzed using SPSS (version 20.0; IBM Corp., Armonk, NY, USA) statistical software. Statistical significance was determined by a p-value<0.05.

Results

A total of 78 patients were included in this study. Among them, 13 had temporal bone fractures and 65 had no temporal bone fractures. The clinical characteristics of the included patients are summarized in Table 1. The mean age of patients, without and with temporal bone fractures, was 53.1 (±14.0) years and 50.1 (±21.9) years, respectively, without any significant difference between the two groups (p=0.788). The mean number of CRPs performed for the remission of BPPV in patients with temporal bone fracture was 2.7 (±1.2), which was larger than that in patients without temporal bone fracture of 2.1 (±0.8). This difference was significant (p=0.045). There was no significant difference in the sex ratios, prevalence of hypertension, diabetes mellitus, dyslipidemia or osteopenia, BPPV side, and type of the affected SCC between the two groups (p>0.05).

The result of univariable analysis showed that temporal bone fracture (p=0.017, β=0.631, 95% CI: 0.118–1.143), diabetes mellitus (p=0.065, β=0.433, 95% CI: -0.027–0.894), and BPPV side (p=0.182, β=-0.266, 95% CI: -0.658–0.127) showed a relatively significant association with the number of CRPs (p<0.2) (Table 2). Multivariable analysis was performed with variables having a p-value of less than 0.2 in the univariable analysis. Only temporal bone fracture showed a significant association with the number of CRPs (p=0.043, β=0.532, 95% CI: 0.017–1.046). The BPPV side and diabetes mellitus showed no statistical significance with the number of CRPs (Table 3).

### Table 1. Demographics of the study population

| Characteristic               | Total | Temporal bone fracture (-) | Temporal bone fracture (+) |
|------------------------------|-------|----------------------------|---------------------------|
| Age (years)                  |       |                            |                           |
| 10–19                        | 3 (3.8)| 2 (3.1)                    | 1 (7.7)                   |
| 20–29                        | 3 (3.8)| 1 (1.5)                    | 2 (15.4)                  |
| 30–49                        | 12 (15.4)| 10 (15.4)             | 2 (15.4)                  |
| 40–49                        | 11 (14.1)| 10 (15.4)               | 1 (7.7)                   |
| 50–59                        | 21 (26.9)| 19 (29.2)               | 2 (15.4)                  |
| 60–69                        | 19 (24.4)| 17 (26.2)               | 2 (15.4)                  |
| 70–79                        | 8 (10.3) | 6 (9.2)                   | 2 (15.4)                  |
| 80–89                        | 1 (1.3) | 0 (0.0)                   | 1 (7.7)                   |
| Mean                         | 52.6 (±15.5)| 53.1 (±14.0)       | 50.1 (±21.9)              |
| Sex                          |       |                            |                           |
| Male                         | 24 (30.8)| 17 (26.2)               | 7 (53.8)                  |
| Female                       | 54 (69.2)| 48 (73.8)               | 6 (46.2)                  |
| Number of canalith repositioning procedure (±SD) | | | |
| Hypertension                 |       |                            |                           |
| Yes                          | 25 (32.1)| 22 (33.8)               | 3 (23.1)                  |
| No                           | 53 (67.9)| 43 (66.2)               | 10 (76.9)                 |
| Diabetes mellitus            |       |                            |                           |
| Yes                          | 18 (23.1)| 13 (20.0)               | 5 (38.5)                  |
| No                           | 60 (76.9)| 52 (80.0)               | 8 (61.5)                  |
| Dyslipidemia                 |       |                            |                           |
| Yes                          | 11 (14.1)| 10 (15.4)               | 1 (7.7)                   |
| No                           | 67 (85.9)| 55 (84.6)               | 12 (92.3)                 |
| Osteopenia                   |       |                            |                           |
| Yes                          | 8 (10.3) | 7 (10.8)                 | 1 (7.7)                   |
| No                           | 70 (89.7)| 58 (89.2)               | 12 (92.3)                 |
| BPPV side                    |       |                            |                           |
| Right                        | 41 (52.6)| 33 (50.8)               | 8 (61.5)                  |
| Left                         | 37 (47.4)| 32 (49.2)               | 5 (38.5)                  |
| Affected canal               |       |                            |                           |
| Posterior                    | 56 (71.8)| 47 (72.3)               | 9 (69.2)                  |
| Lateral                      | 22 (28.2)| 18 (27.7)               | 4 (30.8)                  |

Data are presented as n (%). SD, standard deviation
Discussion

The pathophysiology underlying otoconia detachment in most BPPV cases is unknown, and external shock/traumas, such as temporal bone fractures are estimated to influence the prognosis of BPPV. Therefore, based on the hypothesis that BPPV associated with temporal bone fracture would be more resistant to CRPs, this study retrospectively investigated the factors associated with the number of CRPs performed for the remission of BPPV in patients with and without temporal bone fractures.

According to the results of this study, the mean number of CRPs performed in patients with temporal bone fracture was 2.7, which was larger than that in patients without temporal bone fracture (2.1). The results of multivariable analysis also revealed that BPPV accompanied by temporal bone fracture requires more frequent CRPs than idiopathic BPPV. This finding is similar to that reported in previous studies, which stated that BPPV caused by trauma has a lower treatment success rate than idiopathic BPPV.\textsuperscript{14,15} This interaction is presumed to be a result of compounded damage to the otolith organs in the case of BPPV accompanied by temporal bone fracture. If the otolith organs were severely damaged, the structures supporting the otoconia in the otolith organ might be destroyed.\textsuperscript{10} Even after the successful CRPs, the otoconia might detach from the otolith organs again, leading to persistence of BPPV.

Since the otoconia causing BPPV originates from the otolith organ, the state of the otolith organ should be considered when predicting the prognosis of BPPV. Several studies have evaluated the state of otolith organs using cervical/ocular vestibular evoked myogenic potential (c/oVEMP). Based on these results, they have reported the association between the state of otolith organs and BPPV.\textsuperscript{8,17-23} These studies compared

Table 2. Univariable regression analysis of factors potentially associated with the number of canalith repositioning procedures performed

| Factor                              | p-value | β    | 95% CI          |
|-------------------------------------|---------|------|-----------------|
| Sex                                 |         |      |                 |
| Male                                | 0.403   | -0.181 | -0.608−0.247   |
| Female                              |         |      |                 |
| Age                                 | 0.748   | 0.002 | -0.011−0.015    |
| Temporal bone fracture              |         |      |                 |
| No                                  | Reference |      |                 |
| Yes                                 | 0.017   | 0.631 | 0.118−1.143     |
| Hypertension                        |         |      |                 |
| No                                  | Reference |      |                 |
| Yes                                 | 0.819   | 0.049 | -0.376−0.474    |
| Diabetes mellitus                   |         |      |                 |
| No                                  | Reference |      |                 |
| Yes                                 | 0.065   | 0.433 | -0.027−0.894    |
| Dyslipidemia                        |         |      |                 |
| No                                  | Reference |      |                 |
| Yes                                 | 0.295   | -0.300 | -0.866−0.266    |
| Osteoporosis or osteopenia          |         |      |                 |
| No                                  | Reference |      |                 |
| Yes                                 | 0.888   | -0.046 | -0.700−0.607    |
| BPPV side                           |         |      |                 |
| Right                               | Reference |      |                 |
| Left                                | 0.182   | -0.266 | -0.658−0.127    |
| Affected SCC                        |         |      |                 |
| Posterior canal                     | Reference |      |                 |
| Lateral canal                       | 0.294   | -0.232 | -0.670−0.206    |

CI, confidence interval; BPPV, benign paroxysmal positional vertigo; SCC, semicircular canal

Table 3. Multivariate regression analysis of factors associated with the number of canalith repositioning procedures performed

| Factor                              | p-value | β    | 95% CI          |
|-------------------------------------|---------|------|-----------------|
| Temporal bone fracture              |         |      |                 |
| No                                  | Reference |      |                 |
| Yes                                 | 0.043   | 0.532 | 0.017−1.046     |
| Diabetes mellitus                   |         |      |                 |
| No                                  | Reference |      |                 |
| Yes                                 | 0.097   | 0.384 | -0.071−0.840    |
| BPPV side                           |         |      |                 |
| Right                               | Reference |      |                 |
| Left                                | 0.173   | -0.263 | -0.643−0.118    |

CI, confidence interval; BPPV, benign paroxysmal positional vertigo

Fig. 2. The number of CRPs performed until the remission of benign paroxysmal positional vertigo in the patients with or without temporal bone fracture. CRP, canalith repositioning procedure.
c/oVEMP results between BPPV patients and normal subjects, or between affected and unaffected sides in BPPV patients. As a result, it was reported that abnormal results of c/oVEMP were more commonly measured in BPPV patients or on the affected side. Furthermore, some studies have predicted the prognosis of BPPV based on the state of the otolith organs using c/oVEMP. Lee et al. reported that c/oVEMP response was more often absent in the recurrent BPPV patients. In our previous study using cVEMP and number of BPPV attacks. In conclusion, this study revealed that BPPV accompanied by temporal bone fracture had more severe damage to the otolith organs. However, the condition of otolith organs was not directly measured. In follow-up studies, it would be helpful to measure the condition of otolith organs in BPPV patients with temporal bone fracture and idiopathic BPPV patients using c/oVEMP. In order to consider the number of CRPs performed until the remission of BPPV as resistance to treatment, the interval between CRPs must be constant. As our hospital treats most BPPV patients every 3 to 4 days, the interval between CRPs for BPPV is mostly 3 or 4 days. To keep this more strictly, patients with an interval between CRPs for BPPV other than 3 or 4 days were excluded.

This study has limitations. First, we assumed that patients with BPPV accompanied by temporal bone fracture had more severe damage to the otolith organs. However, the condition of otolith organs was not directly measured. In follow-up studies, it would be helpful to measure the condition of otolith organs in BPPV patients with temporal bone fracture and idiopathic BPPV patients using c/oVEMP. Second, the number of patients with temporal bone fracture included in the study is not large. Future studies will include more patients to obtain more robust results.

In conclusion, this study revealed that BPPV accompanied by temporal bone fracture, inflicting significant damage to the otolith organ, may be more resistant to CRPs than idiopathic BPPV. Therefore, patients with temporal bone fracture warrant more intensive examination and treatment of BPPV than those without temporal bone fracture. In addition, this study contributed to clarifying the pathophysiology underlying the development of BPPV.

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Author Contribution

Conceptualization: all authors. Data curation: Munyoung Chang. Formal analysis: Munyoung Chang, Seog-Kyun Mun. Methodology: Munyoung Chang. Project administration: Munyoung Chang, Seog-Kyun Mun. Visualization: Munyoung Chang, Seog-Kyun Mun. Writing—original draft: Munyoung Chang. Writing—review & editing: Sei Young Lee, Seog-Kyun Mun.

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