

# Vestibular Rehabilitation to the Patients With Vestibular Migraine and Vestibular Neuritis

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## Research

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# Abstract

## Background

It is not yet clarified the effect of vestibular rehabilitation on patients with vestibular migraine (VM) .

## Aims/Objectives

In this study, we aimed to compare the effect of vestibular rehabilitation on patients with VM and those in the chronic stage of vestibular neuritis (VN).

## Material and Methods

A total of 26 patients with VM and 31 patients in the chronic stage of VN who were treated. All patients underwent an in-hospitalized vestibular rehabilitation program. A variety of data including the Dizziness Handicap Inventory (DHI), POMS, and posturography were compared.

## Results

Before treatment, there was no significant difference in the parameters between the two groups, except for a higher confusion score of Profile of Mood States (POMS) in the VM group. In both groups, the DHI score significantly improved. In the VM group, the confusion score of POMS, physical component score (PCS) of SF-8, and some parameters in posturography significantly improved. In the VN group, anxiety, depression, and PCS of SF-8 significantly improved.

## Conclusions and Significance

The higher confusion score of POMS in patients with VM reflects the known increased comorbidity risk of psychiatric disorders. In both groups, in-hospitalized vestibular rehabilitation was effective in alleviating the subjective and objective parameters of vestibular symptoms within a month.

## Introduction

An association between vestibular symptoms, including dizziness and vertigo, and headache, has been reported in several studies [1]. The diagnostic criteria for “vestibular migraine” (VM) are described in the International Classification of Headache Disorders (ICHD), third edition, beta version[2]. Patients with chronic dizziness often complain of not only physical complaints but also mental issues. There are reports on the high rate of psychiatric comorbidity in patients with chronic dizziness, including VM [3, 4], which may contribute to the treatment outcomes.

There is moderate to strong evidence that vestibular rehabilitation is safe and effective for managing unilateral peripheral vestibular dysfunction, based on several high-quality randomized controlled trials [5]. However, there is insufficient evidence for vestibular rehabilitation in patients with VM. Sugaya et al. reported the effect of vestibular rehabilitation in a patient with VM [6]. However, in their study, the effect of rehabilitation was not compared with a specific group of patients with dizziness but was compared with patients with tension-type headache and various states of dizziness. It is clinically useful to compare the effect of vestibular rehabilitation between VM and other vestibular diseases, such as vestibular neuritis (VN), where the efficacy of vestibular rehabilitation has been sufficiently demonstrated [7]. For analysis of the patient’s mood, they only evaluated anxiety and depression using the Hospital Anxiety and Depression Scale. Headache is reported to be associated with various emotions [8]; thus, multidimensional emotions in VM should be investigated. Additionally, the health-related quality of life (QOL) is reportedly lower in participants with VM than in controls without dizziness [9]. Improvement of the health-related QOL in VM is of clinical importance.

This study aimed to compare the characteristics and effects of vestibular rehabilitation on patients with VM and VN. We tried to analyze not only the dizziness handicap but also multidimensional emotions and the mental and physical health-related QOL in the patients.

## Materials And Methods

Patients with a chief complaint of dizziness who visited the Department of Otorhinolaryngology at the National Tokyo Medical Center were included in this study. A retrospective chart review was conducted. A total of 26 patients (4 males, 22 females; average age  $51.2 \pm 18.6$  years) with VM and 31 patients (10 males, 21 females,  $57.1 \pm 12.4$  years) in the chronic stage of VN were treated at our institute from 2015 to 2017. Participants complained of persistent dizziness even after conventional treatment, which included (1) drug therapy with betahistine 36 mg daily for the initial 2–4 weeks, (2) lifestyle counseling regarding exercise and daily walking, and (3) attaining adequate sleep and stress reduction. We recruited participants who met the above-mentioned criteria if (1) the patient was  $\geq 20$  years of age, (2) the dizziness had persisted for at least 3 months despite conventional therapy in the outpatient clinic, (3) the patient wished to have intensive, inpatient therapy for the persistent dizziness, and (4) the patient was literate. The exclusion criteria were as follows: (1) a diagnosis of dizziness due to a cerebrovascular disorder, (2) medical contraindications for the necessary head movements during vestibular rehabilitation (e.g., severe cervical disorder), (3) serious comorbidity (for instance, a life-threatening condition, severe cognitive impairment, or severe psychiatric disorder), (4) central nervous system disease, or (5) bilateral vestibular deficit. The diagnosis of VM was based on the ICHD, third edition, beta version [2] and VN was based on the diagnostic criteria of the Japan Equilibrium Research Association [10]. Patients with VN complained of dizziness for at least 3 months after the diagnosis of VN.

This study was approved by the ethical committee of the National Tokyo Medical Center (R12-009) and was performed in accordance with the ethical standards laid down in the 1964 Declaration of Helsinki and its later amendments.

## Measures

### Dizziness Handicap Inventory

The Dizziness Handicap Inventory (DHI) [11] is a standard questionnaire that quantitatively evaluates the degree of handicap in the daily lives of patients with vestibular disorders and consists of 25 questions. The total score ranges from 0 (no disability) to 100 (severe disability).

### State-Trait Anxiety Inventory

The State-Trait Anxiety Inventory (STAI) [12] is a self-reported questionnaire consisting of 40 questions, comprising a state anxiety subscale with 20 items and a trait anxiety subscale with 20 items. Each item of the STAI assesses how respondents “generally” feel regarding each statement (e.g., “I feel at ease”) on a scale of 1 (almost never) to 4 (almost always).

### Self-Rating Depression Scale

The Self-Rating Depression Scale (SDS), designed by Zung in 1965 [13], is generally considered a reliable instrument for measuring depressive symptoms in primary care. It is a self-reported, 20-item questionnaire. Item responses are rated from 1 to 4, with higher scores corresponding to more frequent symptoms.

### Profile of Mood States (POMS)

The POMS is a psychological rating scale used to assess transient, distinct mood states. The POMS measures six different dimensions of mood swings over a period of time. These include tension or anxiety, anger or hostility, vigor or activity, fatigue or inertia, depression or dejection, and confusion or bewilderment. A five-point scale ranging from “not at all” to “extremely” is administered by experimenters to patients for assessing their mood state.

### Japanese version of the Medical Outcomes Study 8-items Short Form Health Survey

Health-related QOL was evaluated using the Japanese version of the Medical Outcomes Study 8-items Short Form Health Survey (SF-8) questionnaire, the validity and reliability of which have already been confirmed [14]. SF-8 comprises eight items — physical functioning, role limitation due to physical problems, body pain, general health, vitality, social functioning, role limitation due to emotional problems, and mental health. The physical health component summary score (PCS) and mental health component summary score (MCS) were measured using the norm-based scoring method, which is based on a large-scale population study conducted in Japan [15]. Higher scores on these subscales indicate better health-related QOL.

### Measurement of the gravity center fluctuation

The measurement of the gravity center fluctuation for objective assessment of the dizziness severity was performed using a stabilometer (G-5000, Anima Corp., Tokyo, Japan), and the total length of path (LNG) and environmental area (ENV) during eye-opening/closing.

## Intervention

Patients were hospitalized for 5 days in groups of 8–10 individuals; the groups were then taught how to self-perform the 30-min vestibular rehabilitation program [16], which comprised repeatedly performing the vestibulo-ocular reflex (VOR) and vestibulo-spinal reflex (VSR). The VOR training included seven exercises: (1) quick horizontal eye movement, (2) quick vertical eye movement, (3) eye tracking the horizontal direction, (4) eye tracking the vertical direction, (5) horizontal head shaking with gazing at a fixed target, (6) vertical head shaking with gazing at a fixed target, and (7) oblique head tilting with gazing at a fixed target. Each eye or head movement was repeated 20 times. The VSR training consisted of eight static and five dynamic exercises. The eight static exercises were: (1) standing up and sitting down with the eyes open, repeated thrice; (2) standing up and sitting down with the eyes closed, repeated thrice; (3) standing with the eyes closed and feet apart for 20 s; (4) standing with the eyes closed and feet together for 20 s; (5) standing in a tandem posture with the right foot in the front for 20 s; (6) standing in a tandem posture with the left foot in the front for 20 s; (7) standing on the right foot for 20 s; and (8) standing on the left foot for 20 s. The five dynamic exercises were: (1) 180° turn to the left, repeated thrice; (2) 180° turn to the right, repeated thrice; (3) walking in a tandem posture for 10 m; (4) walking with horizontal head shaking for 10 m; and (5) walking with vertical head shaking for 10 m. During the training, patients performed these exercises thrice a day under the supervision of a physician. After 5 days, all patients learned to perform the exercises. The patients were then instructed to continue performing the vestibular rehabilitation program thrice a day after discharge.

## Procedure

After the participants provided written informed consent, they were evaluated on the day of hospitalization (time 1), as well as a month later (time 2), using the questionnaires mentioned previously. Static posturography was also performed. All drugs that could affect dizziness, including vestibular suppressants, were discontinued soon after the introduction of vestibular rehabilitation.

## Statistical analysis

Data analysis was performed using the SPSS 25.0 software (IBM Corp, NY, USA). A two-way repeated measures analysis of variance (ANOVA) was performed to analyze the effects of group and time on all outcomes. The t-test was used for group comparisons of age. A P-value <0.05 was considered significant.

## Results

### Descriptive results

Forty-nine patients (12 males and 37 females, mean age = 54.1±16.3 years) remained in the analysis. The participants in this study comprised 25 patients with VM and 24 patients with VN. There was no significant difference in age between the groups ( $t[41.6] = 1.3$ ,  $p = 0.21$ ). Some patients did not provide any data regarding STAI responses (valid data:  $N = 48$ ), SDS responses (valid data:  $N = 47$ ), POMS responses (valid data:  $N = 47$ ), SF-8 responses (valid data:  $N = 46$ ), and LNG and ENV during eye opening and closing (valid data:  $N = 46$ ). In cases where the participant fell, measurement of their center of gravity fluctuation was treated as missing data.

#### Two-way ANOVA results

Table 1 shows the comparison between groups and time points for all variables using two-way repeated measures ANOVA. There were significant main effects of group on the trait scale of STAI and the SDS score (VM > VN) and significant main effects of time on the DHI score, state and trait scores of STAI, SDS score, trait-anxiety, depression, anger-hostility, confusion, and total mood disturbance scores of POMS, the PCS and MCS scores of SF-8, LNG during eye closing, and ENV during eye closing (SF-8: time 1 < time 2, other variables: time 1 > time 2). Regarding the confusion score of POMS, there was a significant interaction between group and time. The post hoc test showed that the confusion score at time 1 in the VM group was significantly higher than that in the VN group and that the confusion score at time 2 was significantly lower than that at time 1 only in the VM group.

## Discussion

In this study, we successfully indicated that the group undergoing vestibular rehabilitation showed improvement almost all parameters of dizziness, emotions, and QOL for both VM and VN. The improvement of physical and psychological symptoms in patients with VM supports our previous results [6]. Additionally, our provides new findings on the effect on anger, confusion, or QOL, which were not investigated in a previous study [6]. Migraine or VM is closely associated with psychosocial stress and should be assessed using both physical and psychological measurements.

The mechanism of efficacy of vestibular rehabilitation in patients with VM can be explained by the physiological background of the association between vestibular function and headache. Some studies have proposed that the mechanism underlying vestibular dysfunction related to migraine is a parallel activation of the vestibular and cranial nociceptive pathways [8]. Nociceptive and vestibular afferents with neurochemical similarities, including the expression of serotonin, capsaicin, and purinergic receptors[8], converge in brainstem structures such as the parabrachial nucleus, raphe nuclei, and locus coeruleus, and these structures play an important role in modulating the sensitivity of pain pathways [8]. Thus, vestibular symptoms may have a biologically close association with headaches.

In this study, the “confusion” score, a subscale of the POMS, indicated the most prominent difference between patients with VM and patients with VN regarding the efficacy of vestibular rehabilitation. The migraineurs reported significantly greater scores on the confusion-bewilderment subscale of the POMS [17]. We also found that patients with VM had a higher score of confusion of POMS at baseline. Additionally, the confusion score improved with vestibular rehabilitation only in patients with VM. Anxiety in patients with VM was significantly higher than that in those with VN in STAI but not in POMS. This discrepancy may be due to a similar level of anxiety in VM and VN. Patients with VN had a high level of anxiety [18], comparable with that of patients with VM.

Our study had several limitations. First, the number of patients with both VM and VN was small. In the future, we will conduct a similar study with a larger sample and compare sexes and age groups or apply appropriate regression analyses. Second, the study was not prospective but a retrospective chart review. Third, the study periods were relatively short, being only a month. In the future, we will conduct a prospective study with long-term follow-up. Fourth, this study lacked a control group. The effect of vestibular rehabilitation is worth investigating using a randomized controlled protocol in order to obtain more convincing evidence.

## Conclusions

Vestibular rehabilitation contributed to an improvement in vertigo along with dizziness and psychological factors in both patients with VM and VN.

## Declarations

#### Author Contributions

FG drafted the manuscript. MA collected the data. FG organized the research, and NS analyzed the data and checked the manuscript.

#### Declaration of interest

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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## Tables

Table 1. Comparison of all variables between groups and time points (two-way repeated measures analysis of variance).

	VM (Mean±SD)				VN (Mean±SD)				Main effect of group		Main effect of time		Interactio	
	Time 1		Time 2		Time 1		Time 2		F	p	F	p	F	p
DHI	50.1	±20.5	30.1	±26.0	53.1	±16.3	26.2	±16.7	0.0	0.93	70.2	<0.0001	1.5	0.0
STAI														
State	49.7	±8.7	45.5	±9.8	47.6	±12.4	41.2	±8.3	1.7	0.19	12.5	0.001	0.6	0.0
Trait	52.7	±11.2	47.9	±11.7	44.8	±9.7	39.0	±9.2	10.1	0.003	12.5	0.001	0.1	0.0
SDS	48.6	±11.6	43.8	±10.4	43.5	±7.3	38.5	±8.1	4.4	0.04	14.0	0.001	0.0	0.0
POMS														
Tension-Anxiety	8.5	±5.2	6.0	±4.2	6.8	±3.8	5.0	±3.6	1.5	0.22	11.8	0.001	0.3	0.0
Depression	6.4	±4.3	4.8	±3.6	4.6	±3.5	3.5	±4.0	2.4	0.13	8.2	0.006	0.2	0.0
Anger-Hostility	6.2	±4.8	5.4	±3.9	5.2	±4.9	3.5	±2.7	1.7	0.20	6.5	0.02	0.6	0.0
Vigor	5.1	±4.1	7.1	±5.5	6.7	±3.7	7.4	±4.0	0.7	0.40	3.6	0.07	1.0	0.0
Fatigue	10.3	±6.1	8.4	±5.7	8.4	±5.0	7.5	±5.1	0.9	0.35	3.4	0.07	0.5	0.0
Confusion	8.7	±3.9	5.8	±3.3	6.0	±3.1	5.6	±2.8	3.0	0.09	10.1	0.003	6.5	0.0
TMD	34.2	±22.4	23.3	±19.4	24.2	±18.9	18.4	±16.8	2.1	0.15	10.2	0.003	0.9	0.0
SF-8														
PCS	38.9	±8.2	44.3	±8.0	37.2	±9.1	46.0	±6.8	0.0	0.98	27.0	<0.0001	1.5	0.0
MCS	40.4	±8.4	44.0	±9.4	43.4	±10.2	45.3	±5.6	1.0	0.32	4.2	0.047	0.5	0.0
LNG														
Eye-opening	105.6	±36.3	97.4	±29.3	101.6	±38.9	94.7	±34.8	0.1	0.72	2.5	0.12	0.0	0.0
Eye-closing	151.4	±56.1	120.4	±46.4	176.4	±116.9	137.2	±51.4	1.3	0.25	9.3	0.004	0.1	0.0
ENV														
Eye-opening	8.4	±5.5	6.8	±6.1	6.1	±6.5	4.8	±3.3	2.3	0.14	2.8	0.10	0.0	0.0
Eye-closing	10.9	±7.4	7.3	±5.5	10.5	±12.7	7.1	±5.4	0.0	0.87	6.0	0.02	0.0	0.0

\* VM > VN at time 1 (F = 6.8, p = 0.01); time 1 > time 2 in VM (F = 17.6, p <0.0001)

VM (vestibular migraine)

VN (vestibular neuritis)

DHI=Dizziness Handicap Inventory

STAI (State-Trait Anxiety Inventory)

SDS=Self-Rating Depression Scale)

POMS (Profile of Mood Status)

TMD (total mood disturbance)

SF-8 (short form 8)

PCS (physical component score)

MCS (mental component score)

LNG (the total length of path)

