# cVEMP, oVEMP and Caloric Test Results in Individuals with Meniere's Disease

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Abstract: Objective: The present study aimed to characterize cVEMP, oVEMP and caloric test findings in individuals with Meniere's disease.

*Methods*: 29 participants with Meniere's disease and 29 participants with normal hearing were enrolled for the study. Out of 29 participants in the experimental group, 25 of them had unilateral Meniere's disease and 4 of them had bilateral Meniere's disease (Total 33 ears diagnosed with Meniere's and 25 ears with non-Meniere's disease). All the participants underwent routine audiological, cVEMP, oVEMP and Caloric testing.

*Results*: Out of 33 ears with Meniere's disease, 29 ears had absent responses on cVEMP, 23 ears showed absent oVEMP responses, 27 ears had hypo-activity, five ears showed hyper activity and one ear showed normal response to caloric stimulation. In the contralateral ear (25 ears with non-Meniere's disease) both cVEMP and oVEMP were absent in 5 of the ears, cVEMP was absent and oVEMP was present in 13 ears, cVEMP was present and oVEMP was absent in 1 ear, whereas both cVEMP and oVEMP were present in 6 ears in individuals with Meniere's disease. However, the caloric responses were present normally in the contralateral ear in individuals with Meniere's disease.

*Conclusions*: The combination of cVEMP, oVEMP and caloric test provides valuable information regarding localization of hydrops in individuals with Meniere's disease.

Keywords: cVEMP, oVEMP, Caloric test, Meniere's disease, Contralateral ear.

# INTRODUCTION

Meniere's disease is a complex, multifactorial disorder of the inner ear which is the most common cause of the episodic vertigo along with fluctuating hearing loss [1]. In spite of several investigations, the etiology and pathophysiology of Meniere's disease remain controversial and incompletely understood [1]. In majority of the individuals, Meniere's disease is linked to idiopathic endolymphatic hydrops, which is an abnormal increase in the volume of endolymph in the inner ear [2, 3]. The excessive accumulation of endolymph may be due to altered absorption by the endolymphatic duct and sac [4] or due to increased secretion of the endolymph [5].

The different physiological/electrophysiological tests used to diagnose Meniere's disease includes Electrocohleography (EcochG), Glycerol test, CHAMP (cochlear hydrops analysis by masking paradigm), Caloric tests and Cervical Vestibular evoked myogenic potential (cVEMP). In EcochG test, the SP (summating potential)/AP (Action potential) amplitude as well as area ratio are found to be higher in individuals with Meniere's disease [6-9]. An improvement in hearing threshold was found to be associated with glycerol administration in individuals with Meniere's disease [6, 10-14]. The latency shift of wave V of auditory brainstem response (ABR) for click alone and click plus high pass masking noise is reported to be significantly reduced in individuals with Meniere's disease when compared to non-Meniere's normal hearing individuals [15].

However there are some inherent problems with the above mentioned tests. In EcochG, it has been reported that SP is present in 60% of the total normal hearing individuals [16]. Hence, making a diagnosis based on SP/AP (Action potential) ratio becomes difficult. In CHAMP test, the intensity used to record the responses is 60dB nHL [15] and hence an individual with hearing loss more than mild degree cannot be evaluated using this technique.

cVEMP is another electrophysiology test which has been utilized to assess the integrity of vestibulospinal reflex [17- 19]. Pathway of cVEMP includes the saccular macula, inferior vestibular nerve, the lateral vestibular nucleus, the medial vestibulospinal tract, and the motor neurons of ipsilateral SCM muscle [20]. cVEMP has been used as a part of the test battery for Meniere's disease [21- 23] and there is sufficient amount of evidence accumulated on its sensitivity in identifying individuals with Meniere's disease which ranges from 50% to 67% [24- 26].

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Sinha et al.

Another variant of Vestibular evoked myogenic potentials (VEMP) is the ocular vestibular evoked myogenic potential (oVEMP). oVEMP are likely to be produced by synchronous activity in the extraocular muscles, i.e., myogenic potentials [27]. The neuronal pathway for oVEMP *via* the vestibulo-ocular reflex include, activation of the vestibular nerve, vestibular nucleus, medial longitudinal fasciculus, oculomotor nuclei, ocular nerves and to the contralateral extraocular muscles [28]. oVEMP responses (negative peak at 10 ms and a positive peak around 15 ms) are vestibular in origin and most likely originating from the otolith-ocular pathway [29].

oVEMP responses mainly assess the function of otolith organs and superior vestibular nerve. oVEMP have also been found to very useful in diagnosis of different vestibular disorders such as Meniere's disease, superior canal dehiscence syndrome, BPPV and auditory neuropathy [30- 33].

In the literature, there are few studies which have utilised all the three tests (cVEMP, oVEMP and Calorics) to characterize the findings in individuals with Meniere's disease [33, 30]. However, these studies have mainly discussed the results of the ipsilateral ear and not the contralateral ear. There are other studies which indicate an involvement of the contralateral ear in 31% to 37% of individuals with Meniere's disease based on several other tests [18, 34, 35]. Thus the present study aimed to characterize cVEMP, oVEMP and caloric test findings both in the ipsilateral and contralateral ears in individuals with Meniere's disease. The study also aimed at associating the findings of the three tests in individuals with Meniere's disease.

# METHOD

### **Participants**

Two groups of participants were enrolled in the present study, clinical group and the control group. Clinical group consisted of 29 participants (18 males & 11 females) with definite Meniere's disease in the age range of 18 to 55 years (Mean age = 39.25 years). Out of 29 participants, 25 participants had unilateral Meniere's disease and 4 had bilateral Meniere's disease. The diagnosis of definite Meniere's disease was based on symptoms exhibited by the participants and the guidelines proposed by the American Academy of Otolaryngology Head and Neck Surgery [36]. Further diagnosis was also confirmed the bv an otolaryngologist.

Control group consisted of 29 aged matched individuals (17 males & 12 females). All the participants in the control group had normal hearing sensitivity with no middle ear pathology. Additionally these participants did not have any vestibular symptoms and any history or presence of any other otological disorders.

# Instrumentation and Test Environment

calibrated GSI-61diagnostic А two channel audiometer with TDH - 39 headphones, and B-71 bone vibrator was used for pure tone audiometry. Calibrated GSI TYMPSTAR immitance meter was used for tympanometry and relexometry. Intelligent Hearing system (IHS version 4.3.02) was used for recording auditory brainstem responses (ABR) and air conducted cVEMP. Biologic navigator Pro EP instrument with biologic insert was used for ocular VEMP recording. RMS Medicare ENG instrument was utilised for recording caloric responses. All the audiological tests were conducted in the acoustically treated rooms and noise levels during the testing were within permissible limits [37].

# Procedure

A detailed case history was taken for each participant prior to testing. Puretone Air conduction thresholds were obtained from 250Hz to 8000Hz and bone conduction thresholds were determined from 250Hz to 4000Hz at octave frequencies for all the participants. Immittance audiometry was carried out in both ears using a probe tone frequency of 226 Hz. Tympanomery was done initially and then ipsilateral and contralateral acoustic reflex threshold was measured for 500, 1000, 2000, and 4000 Hz stimuli. Uncomfortable loudness level (UCL) was obtained in both ears for air conducted speech stimuli using ascending method. Followed by this, ABR was recorded for both the ears to rule out any retro cochlear pathology. Two channel ABR recording was done for 100µsec click stimuli at 90 dBnHL with the rarefaction polarity. The repetition rate used was 11.1/sec and 90.1/sec. The responses were filtered between 100 Hz to 3000Hz.

#### Vestibular Evoked Myogenic Potentials

# cVEMP Recording

The participants were instructed to sit straight and turn their head to the opposite side of the ear in which stimulus was presented, so as to activate ipsilateral sternocleidomastoid muscle (SCM), as it gives reliable and greater amplitude. cVEMP was recorded using 500 Hz tone burst presented at a rate of 5.1/sec using rarefaction polarity. 500 Hz tone burst stimuli was used as the 500 Hz tone burst stimulus gives better amplitude of the cVEMP [38]. The stimuli were presented to the test ear at an intensity of 95 dBnHL using ER - 3A insert ear phones. The responses were recorded for 70 msec post stimulus period along with the 10 msec pre-stimulus period. The recorded responses were then amplified (X 5000) and band pass filtered between 30 to 1500 Hz. The responses were averaged totally for 200 stimuli. Visual feedback system available in the instrument was utilized during the recording in order to ensure that the subjects monitored the tonic electromyogram (EMG) activity of the SCM and maintained it between 100% to 200 %(  $50\mu\nu$  to  $100\mu\nu$ ) to obtain optimum responses.

#### oVEMP Recording

oVEMP was recorded for all the participants with upper gaze. Participants were instructed to maintain the same upper gaze throughout the test run. 500 Hz tone burst was presented at a rate of 5.1/sec using rarefaction polarity. The stimuli were presented monaurally at an intensity of 95 dBnHL to the contralateral ear using ER-3A insert ear phones. 200 stimuli were used for response averaging. The response was analysed for 60 msec post stimulus period. A pre-stimulus period of 10 msec was utilised to record background electrical activity. The recorded electrical responses were amplified (X 5000) and band pass filtered between 1 Hz to 1000 Hz. oVEMP responses were recorded twice in each ear to ensure replicability of the responses.

### **Caloric Testing**

Prior to the testing, ENG equipment was calibrated for each participant using a calibration light bar. All the participants were asked to stop taking anti-vertigo medications 48 hours before the testing. Participants were also asked not to consume alcohol 48 hours before the testing. In caloric test, open loop water irrigation was used to stimulate the horizontal semicircular canal. The temperature selected for warm stimulation was 44° C and temperature for cold stimulation was 30° C. 200ml of fluid was irrigated over a period of 30 secs. The order of irrigation used were, right 44° C, left 44° C, right 30° C, and left 30° C. Recording was done for 3 minutes including the period for which the fluid was irrigated. A rest period of 7 minutes was given between two successive irrigations. The alertness of participants was maintained

throughout the test by giving simple arithmetic problems.

The cumulative frequency was chosen as the parameter to be represented on the butterfly chart. The response waves obtained in the 4 conditions were analysed and the cumulative frequency was calculated. In order to calculate it, the recordings were divided into 10sec intervals. The 3 adjacent intervals having the most number of nystagmus beats, as determined by manual calculations in each 10sec interval, were considered. Thus, the cumulative frequency represented the total number of beats present over a 30sec period. The cumulative frequency response was plotted on the Claussen butterfly chart as reported earlier [39- 41].

# RESULTS

#### **Test Findings in Control Group**

# cVEMP Results

cVEMP responses could be recorded in all the participants in the control group. In cVEMP, the latency of P1 and N1 peaks, amplitude of P1-N1 complex and amplitude asymmetry (between the two ears) was analyzed. Descriptive statistics was done to find out mean and standard deviation for P1 and N1 latencies, amplitude of P1-N1 complex, and inter ear amplitude asymmetry for P1-N1 complex. The descriptive results of latency, amplitude of P1-N1 complex and inter-ear amplitude asymmetry for P1-N1 complex for the control group is shown in Table **1**.

Parameters	Mean	SD
P1 latency (msec)	15.10	1.24
N1 latency (msec)	22.45	2.05
Amplitude of P1-N1 complex (µv)	40.39	12.66

#### Table 1: Mean and Standard Deviation (SD) Values of Latency and Amplitude Measures of cVEMP in Control Group

#### oVEMP Results

Latency of n1, p1 and n2, peak to peak amplitude of n1-p1 and p1-n2 complex, and inter-ear amplitude asymmetry were analyzed in oVEMP. The oVEMP could be recorded in all the participants of the control group. Descriptive statistics was done to find out the mean and standard deviation of latency and amplitude parameters of oVEMP. Mean and standard deviation of

latency of n1, p1 and n2 and amplitude measures are shown in Table **2**.

#### Table 2: Mean and Standard Deviation (S.D) Values of Latency and Amplitude Measures of oVEMP in Control Group

Parameters	Mean	Standard deviation
n1 latency (msec)	11.37	0.96
p1 latency (msec)	16.49	0.90
Amplitude of n1-p1 complex (µv)	9.02	6.31
Amplitude of p1-n2 complex (µv)	8.16	5.62

#### **Caloric Test**

Bithermal caloric test was recorded from all the subjects in the control group. The culmination frequency was calculated for all the participants in the control group. In control group, the range of culmination frequency of nystagmus in response to different caloric stimulation is shown in Table **3**.

Table 3: Range of Culmination Frequency/30 Seconds for all Four Caloric Stimulation in Control Group.

Caloric stimulation	Range of culmination frequency per 30 seconds
Right warm	22 – 59
Left warm	20 – 70
Right cold	21 – 51
Left cold	22 - 64

Claussen's butterfly chart was made from the culmination frequency obtained from the participants in the control group. Figure **1** shows a butterfly chart obtained from one of the participants in the control group.

# Vestibular Findings in Individuals with Meniere's Disease

A total of 29 subjects in the Meniere's disease group were evaluated using the vestibular evoked myogenic potentials (cVEMP and oVEMP) and caloric test. Out of 29 subjects 4 had bilateral pathology whereas 25 had unilateral pathology.

# cVEMP Results

cVEMP recordings showed that, out of 33 ears (29 participants) with Meniere's disease, 29 ears had



**Figure 1:** Results of caloric test for one participant in control group as shown in butterfly chart.

absence of cVEMP responses (87.87%), and remaining four participants the cVEMP recordings were present. Descriptive statistics was done to find out the mean and standard deviations for P1 latency, N1 latency and P1-N1 amplitude for the four subjects in whom the responses were present. The mean latency of P1 peak was 15.16 msec (SD=1.23 msec), the mean latency of N1 peak was 22.68 msec (SD=2.09 msec) and mean amplitude of P1-N1 peak complex was 40.30 uv (SD=12.90). We did not calculate the inter-ear amplitude asymmetry ratio, as in most of the individuals the cVEMP responses were absent. A Kruskal Wallis test was done to find out the significant difference between normal hearing participants and participants with Meniere's disease. Kruskal Wallis test revealed no significant difference for P1 latency between the two groups (p>0.05), P1-N1 amplitude complex between the two groups (p>0.050, however Kruskal Wallis test showed s significant difference for N1 peak latency between the two groups (p<0.05). cVEMP recording of one participant with Meniere's disease is shown in Figure 2.

# oVEMP Results

In oVEMP recordings, out of 33 ears, 23 ears showed absent responses (69.69%) and 10 ears showed present oVEMP response (30.30%). Descriptive statistics was done to find out the mean and standard deviations for the oVEMP parameters for the Meniere's disease participants for whom the responses were present.



Figure 2: cVEMP responses of one participant with right Meniere's disease.



Figure 3: Ovemp response from one of the participants with Meniere's disease. Figure shows presence of oVEMP in left ear and absence of oVEMP in right ear.

The mean latency for n1 peak was 11.50 msec (SD= 0.93), mean latency for p1 peak was 16.57 msec (SD=0.90), mean latency of N2 peak was 21.71 msec (SD=1.67), the mean amplitude of n1-p1 peak complex was 7.86 µv (SD=5.90), whereas the mean amplitude of p1-n2 complex was 7.09 µv (SD= 5.26). Kruskal wallis test was done to determine the significant difference between normal hearing individuals and individuals with Meniere's disease. Kruskal wallis test did not reveal any significant difference for n1 latency, p1 latency and n2 latency between normal and Meniere's disease individuals (p>0.05), whereas Kruskal wallis test revealed a significant difference for n1-p1 amplitude complex (p<0.05) and also for p1-n2 amplitude complex (p<0.05). oVEMP recording of one participant with right Meniere's disease is shown in Figure 3.

#### **Caloric Test Findings**

In caloric test, out of 33 ears with Meniere's disease, hypo activity was noted in 27 affected ear (81.81%), 5 ears showed hyper activity (15.15%) and one ear showed normal response (3.03%) to caloric stimulation. Figure **4** shows butterfly charts of a hypoactive, hyperactive and a normal response.

# Association of Caloric Test, cVEMP and oVEMP Results in Affected Ears of Subjects with Meniere's Disease

Out of 33 ears, 22 ears showed (66.67%) abnormal results in both caloric test and oVEMP. None of the ear showed normal findings in both the tests. In 20 ears (60.60%) with Meniere's disease, both oVEMP and cVEMP were absent. One ear (3.03%) showed normal



Figure 4: Butterfly chart showing a hypoactive, a hyperactive and a normal response in individuals with Meniere's disease.

response in both the tests. 28 ears showed (84.84%) abnormal results in both caloric test and cVEMP. None of the ear showed normal findings in both the tests.

To find out any significant association between the Calorics tests and cVEMP, Caloric tests and oVEMP, oVEMP and cVEMP responses, a Chi square test was done. Chi-square test revealed no significant association between any of the two tests. Results of chi-square test are shown in Table **4**.

It is evident from the Table **4** that none of the pair of vestibular tests show a significant association between them (p>0.01).

#### Vestibular Findings in Contralateral Ears of Individuals with Meniere's Disease

The intact ears (n=25) of individuals with Meniere's disease, all ears showed normal caloric results. Out of

25 ears, 16 ears had absent responses for cVEMP and 6 ears had absence of responses in oVEMP. Caloric test indicated normal responses in all the contralateral ears considered for the current study. Descriptive statistics was done to find out the mean and standard deviations for the latency and amplitude parameters of the cVEMP and oVEMP and are given in Table **5**.

Kruskal Wallis test was done to find out the significant difference between the cVEMP and oVEMP of Meniere's disease and normal hearing individuals. Kruskal Wallis test results revealed a significant difference for n1 latency of oVEMP (p<0.05) and p1n2 amplitude of the oVEMP between normal and Meniere's disease individuals. For rest of the cVEMP and oVEMP parameters, Kruskal Wallis test did not reveal any significant difference between the two groups.

Table 4:	Association	of Caloric Test	, oVEMP	and cVEMP	in Individuals	with	Meniere's Disease
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Test		oVEMP		cVEMP			
		Normal	Absent	Total	Normal	Absent	Total
Caloric test	Normal	0	1	1	0	4	4
Caloric test	Absent	10	22	32	1	28	29
	Total	10	23	33	1	32	33
	p value*	0.697			0.879		
cVEMP	Normal	1	3	4			
	Absent	9	20	29			
	Total	10	23	33			
	p value*	0.649					

\*Chi-square test.

Table 5: Mean and Standard Deviation (SD) for the<br/>Latency and Amplitude Parameters of the<br/>cVEMP and oVEMP in the Contralateral Ear

Potentials	Parameters	Mean	SD
cVEMP	P1 Latency (msec)	14.89	1.09
	N1 Latency (msec)	22.14	2.29
	P1-N1 amplitude complex (µv)	39.66	13.01
oVEMP	n1 latency (msec)	12.04	0.75
	p1 latency (msec)	16.39	0.86
	n1-p1 amplitude complex (µv)	5.21	2.39
	p1-n2 amplitude complex (µv)	4.95	2.94

# Association of Caloric Test, cVEMP and oVEMP Results in Contralateral Ears of Subjects with Meniere's Disease

To find out any association between the cVEMP, oVEMP and Caloric test in contralateral ears of Meniere's disease a Chi-square test was administered. Chi square test could be administered only between cVEMP and oVEMP and could not be done between caloric test and cVEMP or oVEMP as the responses for caloric test was present normally in all the subject. Results of the Chi-square test are given in Table **6**.

Table 6: Association of cVEMP and oVEMP Results in Contralateral Ears of Individuals with Meniere's Disease

Test		cVEMP				
		Absent	Present	Total		
oVEMP	Absent	5	1	6		
	Present	13	6	19		
	Total	18	7	25		
	p value*	0.50				

\*Chi-square test.

It can be seen in Table **6** that both cVEMP and oVEMP were absent in 5 of the ears, cVEMP was absent and oVEMP was present in 13 of the ears, cVEMP was present and oVEMP was absent in 1 ear, whereas both cVEMP and oVEMP was present in 6 individuals with Meniere's disease. Chi-square test failed to reveal any significant association between cVEMP and oVEMP results in unaffected ears of individuals with Meniere's disease.

# DISCUSSION

In the present study, cVEMP responses were absent in 87.87 % of the participants with Meniere's

disease whereas, in 69.69% individuals with Meniere's disease the oVEMP responses were absent. In the literature there have been equivocal findings regarding the presence or absence of cVEMP in participants with Meniere's disease. De Waele et al. [25] have reported a 54% positive rates on cVEMP Murofushi et al. [17] reported 65% positive rate whereas, Young and groups [42] reported 88% detection rates in cVEMP for individuals with Meniere's disease. Huang et al. [43] reported absence of oVEMP in 44% of the Meniere's disease, whereas, Murofushi et al. [33] reported absence of oVEMP in 50% of the cases with Meniere's disease. Also, Chivarovano et al. [30] reported absence of oVEMP in 70% of the cases with Meniere's disease. The differences in different study might be due to the different stage of Meniere's disease. In the early Meniere's disease the VEMP might be present but might disappear at a later stage [42]. All the participants in the present study were diagnosed as having definite Meniere's disease. We assume that all the participants of this study were in the later stage of Meniere's disease.

In this study, the percentage of absence of cervical VEMP was more compared to the ocular VEMP in individuals with Meniere's disease. In individuals with Meniere's disease, during the latent period, ocular VEMP and cervical VEMP could be affected. Katayama *et al.* [44] showed with previous MRI data that endolymphatic hydrops distension in the vestibule has a large effect on the cervical VEMP. Here, the present data also showed that cervical VEMP were more frequently absent than occular VEMP, suggesting that sacular function could be more affected than the utricular function in individuals with Meniere's diseases.

None of the subject with Meniere's disease showed prolonged latency of either cVEMP or oVEMP. Only latency of N1 peak of the cervical VEMP was prolonged for normal hearing individuals compared to the individuals with Meniere's disease. It has been reported that the latency is not affected by saccular or utricular pathology as a result of Meniere's disease as changes in latency are thought to arise from changes in the neural conduction pathways of the sacculo-collic reflex pathway for the cervical VEMP [23] or utricular ocular pathway for oVEMP [30]. However, a neural delay at the level of the receptor organ may contribute to changes in response latency. Studies by Young et al. [42], Murofushi et al. [17] and Ochi et al. [45] have confirmed this and determined that VEMP latency measures are stable in Meniere's disease. Evidence from the cVEMP & oVEMP data in this study supports

this theory. The prolongation of N1 peak alone could just be a chance factor.

Hypo activity to caloric stimulation in the affected ear was the most common finding in the present study, which is similar to the studies reported in the literature [46]. In the present study, 27 out of 33 affected ears (81.81%) showed hypo activity in caloric test. Hypo activity in caloric response could be due to the damage to the hair cells in the horizontal semicircular canal [33]. Five out of 33 ears diagnosed with Meniere's disease had hyperactive responses. The hyperactive caloric responses in patients who suffer from Meniere's disease may be a transient phenomenon, caused by fluctuations of the vestibular condition, central compensation, age and/or mental state of the patients [47].

Other significant results were obtained in the contralateral ears of the individuals with Meniere's disease. In the contralateral ears (n=25) of the individuals with Meniere's disease cVEMP and oVEMP were absent in 5 of the ears, cVEMP was absent and oVEMP was present in 13 of the ears, cVEMP was present and oVEMP was absent in 1 ear, whereas both cVEMP and oVEMP was present in 6 individuals with Meniere's disease. However, the caloric test showed a normal test results in the contralateral ears of individuals with Meniere's disease. Studies in the literature showed that second ear involvement in individuals with unilateral Meniere's disease was seen in 31% to 37% of cases [34, 35]. Study by Lin et al. [18] found that 27% of participants with unilateral Meniere's disease showed abnormal cVEMP responses in the contralateral ear, which is similar to the present study [18].

Histopathological studies of temporal bones of individuals with Meniere's disease showed that hydrops were more common in saccule and utricle compared to the semicircular canal [48] so it can be concluded that, abnormal cVEMP or oVEMP responses may precede the symptoms in the contralateral ear, so VEMP (cVEMP and oVEMP) responses can be used to predict the chances of involvement of contralateral ear in individuals with Meniere's disease.

In the present study, no statistically significant association between caloric test, cVEMP and oVEMP could be obtained. Significant association only between caloric and oVEMP results, not between caloric and cVEMP or between oVEMP and cVEMP have been reported in peripheral vestibular disorder [33]. It can be hypothesised that the three tests assesses different pathways and extent to which these pathways are affected might vary and hence there might not be any association between the three test results. However, combining caloric test, oVEMP and cVEMP may provide localization of site of lesion in the vestibular labyrinth in affected as well as unaffected ears.

# CONCLUSIONS

Caloric test, cVEMP and oVEMP mainly assess the functioning of semi circular canal ocular reflex pathway, sacculo-collic pathway and utriculo-ocular reflex pathway respectively. In vestibular dysfunction, one or more reflex pathways are affected. The combination of cVEMP, oVEMP and caloric test, provides valuable information regarding the involvement of saccular, utricular and semicircular canal involvement in individuals with Meniere's disease. Also, the cVEMP and oVEMP can provide useful information about the involvement of unaffected ears in individuals with Meniere's disease.

# **CONFLICT OF INTEREST**

The authors report no conflict of interests.

# ACKNOWLEDGEMENT

This study is a part of the Project of AIISH research fund. Financial support for this study has been received from AIISH research Fund. The authors thank the Director, AIISH, for granting the project. The authors also thank all the participants for giving their valuable time for our data collection.

### REFERENCES

- Gates G. Meniere's disease review. J Am Acad Audiol 2006; 17: 16-26.
  - http://dx.doi.org/10.3766/jaaa.17.1.3
- [2] Paparella MM, Griebie MS. Bilaterality of Meniere's disease. Acta Otolaryngol 2004; 97: 233-7. <u>http://dx.doi.org/10.3109/00016488409130984</u>
- [3] Merchant SN, Adams JC, Nadol JB. Pathology and pathophysiology of idiopathic sudden sensorineural hearing loss. Otol Neurotol 2005; 26: 151-60. <u>http://dx.doi.org/10.1097/00129492-200503000-00004</u>
- [4] Arenberg IK, Gibson WPR, Bohlen HKH. Improvements in audiometric and electrophysiologic parameters following nondestructive inner ear surgery utilizing a valved shunt for hydrops and Meniere's disease. Proceedings of the Sixth Annual Workhops on Electrocochleography & Otoacoustic Emissions. Denver, CO: International Meniere's Disease Research Institute 1993; 545-61.
- [5] Hallpike CS, Cairns H. Observations on the pathology of Meniere's syndrome. J Laryngol Otol 1938; 53: 625-54. <u>http://dx.doi.org/10.1017/S0022215100003947</u>

- [6] Yen PT, Lin CC, Huang TS. A preliminary report on the correlation of vestibular Meniere's disease with electrocochleographyand glycerol test. Acta Otolaryngol 1995; 115 (520): 241-6. <u>http://dx.doi.org/10.3109/00016489509125238</u>
- [7] Ferraro JA, Durrant JD. Electrocochleography in the evaluation of patients with Meniere's disease/endolymphatic hydrops. J Am Acad Audiol 2006; 17(1): 45-68. http://dx.doi.org/10.3766/jaaa.17.1.6
- [8] Baba A, Takasaki K, Tanaka F, Tsukasaki N, Kumagami H, Takahashi H. Amplitude and area ratios of summating potential/action potential (SP/AP) in Meniere's disease. Acta Otolaryngol 2009; 129(1): 25-29. <u>http://dx.doi.org/10.1080/00016480701724888</u>
- [9] Iseli C, Gibson W. A comparison of three methods of using transtympanic electrocochleography for the diagnosis of Meniere's disease: click summating potential measurements, tone burst summating potential amplitude measurements, and biasing of the summating potential using a low frequency tone. Acta Otolaryngol 2010; 130(1): 95-101. http://dx.doi.org/10.3109/00016480902858899
- [10] Fukuoka H, Takumi Y, Tsukada K, et al. Comparison of the diagnostic value of 3 T MRI after intratympanic injection of GBCA, electrocochleography, and the glycerol test in patients with Meniere's disease. Acta Otolaryngol 2012; 132(2): 141-5. http://dx.doi.org/10.3109/00016489.2011.635383
- [11] Cen J, Zeng X, Wang S, Li Z, Zhang G, Liu X. Analysis of factors affecting the pure tone threshold glycerol test. Lin Chung Er Bi Yan Hou Tou Jing Wai Ke Za Zhi 2010; 24(18): 838-40.
- [12] Mom T, Gilain L, Avan P. Effects of glycerol intake and body tilt on otoacoustic emissions reflect labyrinthine pressure changes in Menière's disease. Hear Res 2009; 250(1-2): 38-45.

http://dx.doi.org/10.1016/j.heares.2009.01.008

- [13] Zhao R, Zhu W, Liu H. The control study of glycerol test in different stage of Meniere's disease patients. Lin Chuang Er Bi Yan Hou Ke Za Zhi 2005; 19(12): 543-4.
- [14] Jablonka A, Pospiech L, Orendorz-Fraczkowska K. Evaluation of glycerol test in Meniere's disease with pure tone audiometry and distortion product otoacoustic emission. Otolaryngol Polish 2003; 57(5): 731-7.
- [15] Don M, Betty K, Tanaka C. A Diagnostic Test for Meniere's Disease and Cochlear Hydrops: Impaired High-Pass Noise Masking of Auditory Brainstem Responses. Otol Neurotol 2005; 26(4): 711-22. http://dx.doi.org/10.1097/01.mao.0000169042.25734.97
- [16] Sass K, Densert B, Arlinger S. Recording techniques for transtympanic electrocochleography in clinical practice. Acta Otolaryngol (Stockh) 1997; 118: 17-25.
- [17] Murofushi T, Shimizu K, Takegoshi H, Cheng PW. Diagnostic value of prolonged latencies in the vestibular evoked myogenic potential. Arch Otolaryngol Head Neck Surg 2001; 127: 1069-72. http://dx.doi.org/10.1001/archotol.127.9.1069
- [18] Lin M, Timmer FCA, Oriel BS, et al. Vestibular evoked myogenic potentials (vemp) can detect asymptomatic saccular hydrops. Laryngoscope 2006; 116: 987- 92. <u>http://dx.doi.org/10.1097/01.mlg.0000216815.75512.03</u>
- [19] Rauch SD. Vestibular evoked myogenic potentials. Curr Opin Otolaryngol Head Neck Surg 2006; 14: 299-304. http://dx.doi.org/10.1097/01.moo.0000244185.65022.01
- [20] Halmagyi GM, Curthoys IS. Clinical testing of otolith function. Ann N Y Acad Sci 1999; 871: 195-204. <u>http://dx.doi.org/10.1111/j.1749-6632.1999.tb09185.x</u>
- [21] Colebatch JG, Halmagyi GM, Skuse NF. Myogenic potentials generated by a click-evoked vestibulocolic reflex. J Neurol Neurosurg Psychiatry 1994; 57: 190-7. <u>http://dx.doi.org/10.1136/jnnp.57.2.190</u>

- [22] Bath AP, Harris N, McEwan J, Yardley MP. Effect of conductive hearing loss on the vestibule-collic reflex. Clin Otolaryngol Allied Sci 1999; 24(3): 181-3. <u>http://dx.doi.org/10.1046/j.1365-2273.1999.00234.x</u>
- [23] Rauch SD, Silveira MB, Zhou G, Kujawa SG, Guinan JJ, Herrmann BS. Vestibular evoked myogenic potentials versus vestibular test battery in patients with Meniere's disease. Otol Neurotol 2004; 25: 981-6. <u>http://dx.doi.org/10.1097/00129492-200411000-00020</u>
- [24] Ribeiro S, De Almeida RR, Caovilla HH, Gananga MM. Vestibular evoked myogenic potentials in the affected and asymptomatic ears in the unilateral Meniere's disease. Braz J Otorhinolaryngol 2006; 71 (1): 60-6.
- [25] De Waele C, Huy PT, Diard JP. Saccular dysfunction in Meniere's disease. Am J Otolaryngol 1999; 20(2): 223-32.
- [26] Kuo SW, Yang TH, Young YH. Change of vestibular evoked myogenic potentials after Meniere's attack. Ann Otol Rhinol & Laryngol 2005; 114 (9): 717-21. <u>http://dx.doi.org/10.1177/000348940511400911</u>
- [27] Rosengren SM, Todd NP, Colebatch JG. Vestibular-evoked extraocular potentials produced by stimulation with boneconducted sound. Clin neurophysiol, 2005; 116: 1938-48. <u>http://dx.doi.org/10.1016/j.clinph.2005.03.019</u>
- [28] Rosengren SM, Welgampola MS, Colebatch JG. Vestibular evoked myogenic potentials: past, present and future. Clin neurophysiol 2010; 121: 636-51. <u>http://dx.doi.org/10.1016/j.clinph.2009.10.016</u>
- [29] Chihara Y, Iwasaki S, Ushio M, Murofushi T. Vestibularevoked extraocular potentials by air-conducted sound: Another clinical test for vestibular dysfunction. Clin neurophysiol 2007; 118: 2745-51. http://dx.doi.org/10.1016/j.clinph.2007.08.005
- [30] Chiarovano E, Zamith F, Vidal PP, De Waele C. Ocular and cervical VEMPs: a study of 74 patients suffering from peripheral vestibular disorders. Clin neurophysiol 2011; 122: 1650-9.

http://dx.doi.org/10.1016/j.clinph.2011.01.006

- [31] Huang CH, Wang SJ, Young YH. Correlation between caloric and ocular vestibular evoked myogenic potential test results. Acta Otolaryngol 2012; 132: 160-6. http://dx.doi.org/10.3109/00016489.2011.624120
- [32] Jacobson GP, McCaslin DL, Piker EG, Gruenwald J, Grantham SL, Tegel L. Patterns of Abnormality in cVEMP, oVEMP, and Caloric tests may provide topological information about vestibular impairment. J Am Acad Audiol 2011; 22: 601-11. http://dx.doi.org/10.3766/jaaa.22.9.5
- [33] Murofushi T, Nakahara N, Yoshimura E, Tsuda Y. Association of air-conducted sound oVEMP findings with cVEMP and caloric test findings in patients with unilateral peripheral vestibular disorders. Acta Otolaryngol 2011; 131 (9): 945-50. http://dx.doi.org/10.3109/00016489.2011.580003
- [34] Thomas K, Harrison MS. Long term follow up of 610 cases of Meniere's disease. Proc R Soc Med 1971; 64(8): 853-6.
- [35] Green JD, Blum DJ, Harner SG. The longitudinal follow up of patients with Meniere's disease. Otolaryngol Head Neck Surg 1991; 104(6): 783-8.
- [36] American Academy of Otolaryngology Head and Neck Surgery Guidelines for the evaluation of therapy in Meniere's disease. Otolaryngol Head Neck Surg 1995; 113, 181-5.
- [37] American National Standards Institute. (1991). American National Standards for maximum permissible ambient noise levels for audiometric test room (ANSI S3.1- 1991). New York: American National Standards Institute.
- [38] Kumar K, Sinha SK, Bharti AK, Barman A. Comparison of Vestibular evoked myogenic potentials elicited by click & short duration one burst. J Laryngol Otol 2011; 125: 343-347. <u>http://dx.doi.org/10.1017/S0022215110001908</u>

- [39] Claussen CF, Von Schlachta I. Butterfly chart for caloric nystagmus evaluation. Archives Otolaryngol 1972; 96: 371-5. http://dx.doi.org/10.1001/archotol.1972.00770090547015
- [40] Kirtane V, Merchant SN, Medikeri SB. Experiences with butterfly chart. J Laryngol Otol 1986; 100: 157-64. <u>http://dx.doi.org/10.1017/S002221510009890X</u>
- [41] Sinha SK, Barman A, Singh NK, Rajeshwari G, Sharanya R. Involvement of peripheral vestibular nerve in individuals with auditory neuropathy. Eur Arch Otorhinolaryngol 2013; 270: 2207-14. http://dx.doi.org/10.1007/s00405-012-2272-4
- [42] Young YH, Huang TW, Cheng PW. Assessing the stage of Meniere's disease using vestibular evoked myogenic potentials. Arch Otolaryngol Head Neck Surg 2003; 129: 815-8. <u>http://dx.doi.org/10.1001/archotol.129.8.815</u>
- [43] Huang CH, Wang SJ, Young YH. Correlation between caloric and ocular vestibular evoked myogenic potential test results. Acta Otolaryngol 2012; 132: 160-6. <u>http://dx.doi.org/10.3109/00016489.2011.624120</u>

Received on 08-06-2015

Accepted on 10-07-2015

Published on 17-09-2015

DOI: http://dx.doi.org/10.12970/2311-1917.2015.03.02.6

- [44] Katayama N, Yamamoto M, Teranishi M, *et al.* Relationship between endolymphatic hydrops and vestibular evoked myogenic potential. Acta Otolaryngol 2010; 130(8): 917-23. http://dx.doi.org/10.3109/00016480903573187
- [45] Ochi K, Ohashi T, Nishino H. Variance of vestibular-evoked myogenic potentials. Laryngoscope 2001; 111: 522-7. http://dx.doi.org/10.1097/00005537-200103000-00025
- [46] Bergman B, Stahle J. Caloric Reaction in Menière's Disease. A Nystagmographic Study of 300 Patients. Acta Oto Laryngol Suppl 1967; 63: 77-9. http://dx.doi.org/10.3109/00016486709123555
- [47] Ikeda M, Watanabe I. Evaluation of hyperactive responses in inner ear diseases. J Otorhinolaryngol Relat Spec 1997; 59(6): 326-31. <u>http://dx.doi.org/10.1159/000276965</u>
- [48] Okuno T, Sando I. Localization, frequency and severity of endolymphatic hydrops and the pathology of the labyrinthine membrane in Meniere's disease. Annals of Otol Rhinol and Laryngol 1987; 96: 438-45. <u>http://dx.doi.org/10.1177/000348948709600418</u>