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Physiology

## A Comparative Study of Brainstem Auditory Evoked Responses in Patients of Type 2 Diabetes Mellitus and Healthy Matched Controls at Tertiary Care Hospital, Bikaner, Rajasthan India

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## **Original Research Article**

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Abstract: Diabetes Mellitus has been implicated as an independent causative factor of sensorineural hearing loss. The Brainstem auditory evoked responses (BAER) represent a useful, non invasive and simple procedure to detect both acoustic nerve and CNS damage. The study was conducted to compare Brainstem auditory evoked responses in patients of Type 2 diabetes mellitus with matched normal subjects. Effect of duration of diabetes on BAER and usefulness of this screening tool were also assessed. A Comparative Cross-sectional study was carried out to compare Brainstem auditory evoked responses in patients of type 2 Diabetes Mellitus (n=50) and matched healthy controls (n=50) at Department of Physiology in association with ENT Department at S.P. Medical College & P.B.M. hospital Bikaner, Rajasthan from June 2016 to May 2017 with Consecutive sampling technique. All BAER waves I to V & Interpeak latencies (I-III, III-V & III-V) of both ears showed statistically significant difference (p<0.05) among cases and controls except Interpeak latency of III-V of left ear. Cases had prolonged interpeak latencies which is an early sign of central neuropathy and can prove diagnostically important. Correlation coefficient among BAER Interpeak latencies among both right and left ears of cases showed only partial or weak correlation which depicts that interpeak latencies are not affected much by duration of disease. Therefore it is derived that BAER waves can act as a screening tool for central neuropathy.

Keywords: Diabetes Mellitus, Brainstem, neuropathy.

### **INTRODUCTION**

Diabetes Mellitus, a chronic multi-systemic metabolic disorder, is fast emerging as an epidemic in both developed and developing countries. The total number of people with diabetes worldwide is projected to rise from 171 million in 2000 to 366 million in 2030. Diabetes Mellitus has been implicated as an independent causative factor of sensorineural hearing loss[1]. Neuropathy, both central and peripheral, is an important complication of Type 2 DM [2]. Neuropathy is the more precocious and frequent late complication of DM. So far most of the clinical and diagnostic studies on diabetic neuropathy have concerned only peripheral and autonomic nerve but recently with the refinement of evoked potential techniques detailed exploration of sensory pathway in central nervous system has been possible. The electrophysiological testing reflects the bioelectric responses of the nervous system to sensory (somatosensory evoked potentials), auditory (brainstem auditory evoked potentials) or

visual stimuli (visual evoked potentials) [3]. The Brainstem auditory evoked responses (BAER) represent a useful, non invasive and simple procedure to detect both acoustic nerve and CNS damage. Brainstem auditory evoked potentials (BAEP) are the potentials that are recorded in response to brief auditory stimulation to assess the conduction through the auditory pathway, up to the midbrain.

## MATERIALS AND METHODS

A Comparative Cross-sectional study was carried out to compare Brainstem auditory evoked response in Patients of type 2 Diabetes Mellitus and healthy controls at Department of Physiology in association with ENT Department at S.P. Medical College & P.B.M. hospital Bikaner, Rajasthan from June 2016 to May 2017 with Consecutive sampling technique. 50 patients of type 2 Diabetes Mellitus patients and 50- healthy controls were assessed. Both cases and controls were matched according to age and sex.

Inclusion Criteria was only proven or diagnosed cases of Type 2 diabetes by history, clinical examination, and blood investigations. The patients conductive deafness, with severe anemia, cardiovascular complications, respiratory diseases, renal failure, patients with past history of neurological disorders, smokers, alcoholic and patients taking any other drugs except for diabetes were excluded. Neurosoft Auditory Evoked Potential (AEP) software was used. Start testing at 70 dB normal Hearing Loss (nHL) and as per response we can increase or decrease the intensity for next recording. Procedure was started by collecting an average of 2000 clicks per intensity. Duration of wave I, II, III, IV and V in milliseconds and Interpeak latencies I-III, I-V and III-V in milliseconds were recorded. The findings were recorded on a predesigned Proforma and statistically analyzed with the help of SPSS 22.0 software.

### RESULTS

In this study 31 (62%) were males and 19 (38%) were females in both groups- diabetics (cases) and control. In both group the age range was of 42-78 years (both groups were matched in terms of age and sex). In control group Mean BMI was  $30.27\pm2.06$ ;

Mean fasting blood sugar was  $72.22\pm2$  mg/dl; Mean HbA1c level was  $4.78\pm1.47$ . Among diabetics Mean BMI was  $39.68\pm4.18$ ; Mean HbA1c  $10.56\pm3.42$  and mean duration of disease was observed to be 10.4 years.

In comparison of waves of BAER in Left ear of both controls & cases (table 1) the difference was statistically significant among all type of waves (p<0.001). and interpeak latencies I-III, I-V showed statistically significant difference (P<0.01) whereas III-V interpeak latency had no statistically significant difference (p>0.05). Comparison of waves & interpeak latencies of Right ear of both controls & cases (table 2) showed that the all BAER waves and all interpeak latencies of statistically significant difference (p<0.05). Correletion coefficient among BAER waves of Left and Right ears with duration of disease among Cases (table 3) shows that only wave II of Right ear is showing moderately positive correlation with duration of disease among cases. All other waves have weak or partial positive correlation (r<0.3). Table 4 shows Correlation coefficient among BAER Interpeak latencies among both right and left ears of cases. This shows only partial or weak correlation which depicts that interpeak latencies are not affected much by duration of disease.

<b>Table-1: Comparison</b>	of waves and interpeal	k latencies of BAER in	Left ear of both controls & cases
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Waves	Cases	Controls	Value	Inference
waves	Mean $\pm$ SD	Mean $\pm$ SD	Of 't'	
I (ms)	$1.59\pm0.30$	$1.25\pm0.15$	7.17	P<0.001
II (ms)	$2.61\pm0.23$	$2.39\pm0.30$	4.11	P<0.001
III (ms)	$4.08\pm0.43$	$3.44\pm0.22$	9.37	P<0.001
IV (ms)	$4.59\pm0.25$	$4.30\pm0.18$	6.66	P<0.001
V (ms)	$6.11\pm0.62$	$5.43\pm0.26$	7.15	P<0.001
I-III (ms)	$2.49\pm0.53$	$2.19\pm0.27$	3.57	P<0.001
I-V (ms)	$4.52\pm0.72$	$4.18\pm0.31$	3.07	P = 0.003
III-V (ms)	$2.03\pm0.66$	$1.98\pm0.36$	0.47	P = 0.64
(Non-Significant: p>0.05, Significant: p<0.05)				

Waves	Cases	Control	Value	Inference
	Mean $\pm$ SD	Mean $\pm$ SD	Of 't'	Interence
I (ms)	$1.55\pm0.33$	$1.18\pm0.15$	7.22	P<0.001
II (ms)	$2.59\pm0.23$	$2.42\pm0.30$	3.18	P= 0.002
III (ms)	$3.58\pm0.43$	$3.45\pm0.21$	1.92	P< 0.05
IV (ms)	$4.43\pm0.21$	$4.30\pm0.20$	3.17	P= 0.002
V (ms)	$6.25\pm0.61$	$5.36\pm0.28$	9.38	P<0.001
I-III (ms)	$2.03\pm0.47$	$2.27\pm0.26$	-3.16	P= 0.002
I-V (ms)	$4.70\pm0.66$	$4.17\pm0.32$	5.11	P<0.001
III-V (ms)	$2.67\pm0.81$	$1.90\pm0.36$	6.14	P<0.001

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Parameters (waves)	Duration of disease ('r') value	Inference
I (ms) Lt	0.082	Weakly positive
II (ms) Lt	0.202	Weakly positive
III (ms) Lt	-0.271	Weakly negative
IV (ms) Lt	-0.21	Weakly negative
V (ms) Lt	0.101	Weakly positive
I (ms) Rt	0.089	Weakly positive
II (ms) Rt	0.341	Moderately positive*
III (ms) Rt	0.049	Weakly positive
IV (ms) Rt	0.078	Weakly positive
V (ms) Rt	0.146	Weakly positive

## Table-3: Correlation coefficient among BAER waves of Left and Right ears with duration of disease among Cases

# Table-4: Correlation coefficient among BAER interpeak latencies (ms) of Left and Right ears with duration of

	disease	
Parameters (interpeak latencies)	Duration of disease ('r') value	Inference
I-III (ms) Lt	-0.26	Weak negative correlation
I-V (ms) Lt	0.05	Very weak positive correlation
III-V (ms) Lt	0.275	Partial positive
I-III (ms) Rt	-0.01	Weak negative
I-V (ms) Rt	0.26	Weakly positive
III-V (ms) Rt	0.22	Weakly positive



Graph-1: Correlation of Wave I of Rt ear with disease duration





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#### DISCUSSION

Different waves of BAER have different source of generators and hence these waves reflect activity of their generators. Wave I represents peripheral nervous system involvement as any change in it depicts the effect of diabetes on auditory nerve. Other absolute waves represent central nervous system involvement as any change in them depicts effect of diabetes on brain stem as generators of waves II, III, IV and V cochlear nucleus, superior olivary nucleus, lateral lemniscuses and inferior colliculus respectively. IPL I-III measures neuronal conduction of acoustic nerve across subarachnoid space into core of lower pons. IPL I-V measures central neuronal conduction from proximal acoustic nerve through pons to midbrain. IPL III-V measures or indirectly reflects neuronal conduction from lower pons to midbrain. It helps in exploring early sub clinical neurological dysfunction in metabolic disorders. BAER might be useful diagnostic tool in detecting central diabetic neuropathy. Controls displayed almost similar values of Mean duration of waves in right and left ears. Brainstem auditory evoked potentials of Left and Right ears among cases show a slight variation among mean values of right and left ear wave duration. Left ear waves have slightly higher peaks than right ear among cases. While comparing BAER waves I-V & Interpeak latencies among cases and controls left ear all five waves duration and interpeak latencies I-III. I-V showed statistically significant difference as cases had prolonged interpeak latencies which is an early sign of central neuropathy and can prove diagnostically important (P<0.001) whereas III-V interpeak latency had no statistically significant difference (p>0.05) which may be explained by observer variation in measurement tool. While all BAER waves and all interpeak latencies of Right ear of both cases and controls showed statistically significant difference (p<0.05) with higher values in right ear of cases showing that BAER may prove a sensitive tool for early diagnosis of neuropathy. Dolu et al. [4], Al-Azzawi et al. [5], Gupta R et al. [6], C Huang et al. [7], observed similar results in their study. Whereas Ologe et al. [8] did BAER on children of age group 9 - 19years with family history of Type 2 diabetes and found no difference of audiogram in children without genetic predisposition. Gupta S. et al. [9] in their crosssectional study observed no significant difference between diabetic and control subjects as regards to the latency of wave IV unilaterally in the left ear and the latencies of waves I, II and interpeak latency I-III bilaterally. Baweja P et al. [10] in their case control study observed significant difference in right ear values but not in left ear values. Shatdal et al. [11] in their cross-sectional study observed that Mean peak latency of waves I, III, V and interpeak latency of I-III, III-V, I-V were prolonged in group 1, but were not statistically significant. Siddiqui SS et al. [12] also observed similar results as ours. Similarly, Mahalik D et al. [13] found that type 2 DM patients showed

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significant prolonged absolute latencies of I, III (P =0.001) and interpeak latencies I-III, III-V and I-V in left ear (P = 0.001) and absolute latencies of I, V (P =0.001), interpeak latencies III-V was statistically significant in right ear. Sushil MI [14], Bhattarai U et al. [15], Anshul sharma et al. [16], Praveen S. Yousuf et al. [17], R.K. Murugeshan et al. [18], Dr K Kanaan [19] observed similar results as our study. Similar to our results, Baweja P et al. [10] in their case control study observed that none of the BAEP latencies were significantly correlated with either the duration of disease or with fasting blood glucose levels in diabetics. Shatdal et al. [11] in their cross-sectional study also observed no significant relation between abnormal BAEP response with age, sex, type of diabetes, duration of diabetes since detection, fasting plasma sugar level, postprandial plasma sugar level, glycosylated haemoglobin, presence of retinopathy, nephropathy and peripheral neuropathy. Whereas Anshul sharma et al. [16] observed a positive correlation between prolongation of latencies and duration of diabetes mellitus. The latencies were also found to be prolonged with altered blood glucose levels.

## CONCLUSION

Therefore it is derived from results that BAER waves as well as interleaf latencies durations are affected among type 2 diabetic patients and can act as a screening tool for central neuropathy whereas duration of diabetes does not prove to have an impact on BAER responses. Hence regular blood glucose monitoring and timely check-up the otorhinolaryngology's and audiologist can resolve many associated problems at an initial stage. Which can certainly improve the quality of life of the individual?

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