Examination and Comparison of Electrically Evoked Compound Action Potentials and Electrically Evoked Auditory Brainstem Response Results of Children with Cochlear Implantation without Inner Ear Anomaly

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Abstract

Objective: To investigate the relationship between electrically evoked compound action potentials (ECAP) and electrically evoked auditory brainstem responses (EABR) in children with cochlear implants (CI) without inner ear anomalies.

Methods: Sixteen children between the ages of two and six years who were CI users participated in the study. ECAP thresholds were recorded from one electrode in the basal, medial, and apical regions of the cochlear implant. EABRs were recorded from electrodes whose ECAP thresholds were determined. The latency-intensity functions, amplitude and morphological analyzes of the eIII and eV waves at 200 and 180 current unit (CU) excitation levels were performed. The data obtained were analyzed statistically.

Results: ECAP thresholds were found to be 171.5±11.38, 169.69±20.32 and 160.81±20.03 CU at the basal, medial and apical electrodes, respectively. EABR thresholds were also found to be 169.69±12.17, 165.62±16.41 and 160.5±15.49 CU in basal, medial and apical electrodes, respectively. There was a strong positive correlation between ECAP and EABR thresholds in apical, medial and basal electrodes (p<0.05).

Conclusion: Consistency was found between ECAP and EABR recordings. However, it was concluded that one could not be preferred over the other because the data quality of the two tests was different. In future studies, ECAP and EABR recordings may be recommended by selecting more electrodes for stimulation.

Keywords: Cochlear implant, electrophysiological studies, electrically evoked compound action potential, electrically evoked auditory brainstem response

Introduction

Cochlear implantation is an amplification method used in children and adults with severe and profound sensorineural hearing loss. Effective programming of speech processors, effectively determining the dynamic range, and ensuring that the patient comfortably perceives acoustic stimulants is highly important in cochlear implant (CI) patients (1).

Electrophysiological measurements are performed to assess and adjust the CI in pediatric patients. These include electrically evoked auditory brainstem response (EABR), electrically evoked stapedial reflex threshold (ESRT), electrically compound action potentials (ECAP), middle latency responses (MLR), and cortical responses (CR). ECAP is the most frequently used of these methods, both intra- and postoperatively. That it provides limited information about auditory nerve fibrils and creates electrical measurement artefact is its disadvantages. On the other hand, it provides detailed information related to the auditory evoked brainstem activity. However, requires deep sleep or sedation because it is affected by muscle artefact (2–4).

The aim of this study is to identify the relationship between the results of two electrophysiological...
techniques, ECAP and EABR, in pediatric CI users with no inner ear anomaly. The hypothesis of the study was defined as "The relationship between ECAP and EABR results are linear."

Methods
This study was conducted at the Dokuz Eylül University Hospital, Otorhinolaryngology Department, Hearing-Speech-Balance Unit, included 16 Nucleus CI 24 Contour users aged from two to six years and were radiologically confirmed to have no abnormalities of the inner ear. Consent was obtained from the families of the patients prior to the study. Ethics committee approval was obtained from the Ethics Committee of the Dokuz Eylül University. ECAP recordings were made while patients were awake. Electrical stimuli were created with apical (electrode number 20), medial (electrode number 11) and basal (electrode number 3) electrodes in ECAP and EABR. Recording electrodes were 22, 13 and 5, respectively. ECAP was measured with 25 microsecond/pulse and 50 Hz stimulus and standard monopolar stimulus (MP1) was used.

Evoked auditory brainstem responses was recorded on a different day than ECAP. Families were asked to keep the child awake the night before and the recording was made during natural sleep using a laptop computer with Medelec Synergy EMG/EP Systems (VIASYS Healthcare, UK) software and equipment. The EABR test protocol was completed in two sessions in some patients. Stimuli were transmitted from the user's processor via a coil by the same CI firm. Connection between the CI and the EABR device was established by an external trigger cable compatible with this brand. Three gold disc electrodes were used in recording and each electrode was carefully placed in the same area during every recording. The difference between positive (forehead-hairline border) and negative (mastoid contralateral to the implanted ear) electrodes was recorded in EABR. The earth electrode was placed in the forehead region (Fpz). Electrodes were placed with skin impedance below 2 kOhms. EABR recording was carried out at 25 Is/phase intervals and using 20 Hz monopolar biphasic alternating polarity stimulation. Responses were filtered using 0.1–3000 Hz analog band-pass filter and amplifier sensitivity was set at 500 µV. Analysis window was 10 msec and sampling rate was 25 kHz. Stimulation was single-channel and simultaneous with the recording. Recording was performed from apical-medial and basal electrodes, respectively. Recording began from the apical region where the most distinct response was evoked (3, 5, 6). Contralateral placement technique was used to minimize artefact (4, 7-10). Intensity-latency function and morphological analysis of the evoked waves were recorded at 200 and 180 current unit (CU) stimulus levels. Before stimulating at these levels, care was taken not to exceed the comfort level set in the most recent programs of the individuals. Following the recordings at 200 CU and 180 CU, stimulus intensity was reduced in increments of 10-units and the last stimulus level at which wave eV was observed was accepted as the EABR threshold. Stimulation levels 5 units above and below the threshold were also recorded (11, 12). At least two tracings were recorded at each stimulation level and the average number in each tracing was set to 1000. ECAP and EABR thresholds of basal, medial and apical electrodes, wave eIII and eV latencies at EABR 200 CU and 180 CU and eIII-eV interwave intervals, amplitude levels of eI and eV waves at this intensity level, and wave eV morphologies were evaluated.

Statistical Analysis
Descriptive statistics of nominal, ordinal and numerical data were performed. The significance of the relationship between ECAP and EABR thresholds was analyzed by Spearman correlation test. Wilcoxon signed-rank test was used to analyze the significance of the difference between ECAP and EABR thresholds. Nonparametric Friedman variance analysis was used to determine whether wave eIII and eV latencies and amplitudes evoked in EABR differed in apical, medial and basal electrodes. A p-value <0.05 was accepted as significant. Statistical Package for the Social Sciences version 16.0 (IBM Corp.; Armonk, NY, USA) software was used.

Results
Of the 16 patients included in the study, 12 were male (75%) and four were female (25%). The CI was in the right ear in 15 patients (94%) and in the left ear in one patient (6%). The mean age of the patients was 46.08±9.13 months and the mean age of the CIs was 27.03±8.2 months. ECAP and EABR responses were evoked by all apical, medial and basal electrodes in all participants. While wave eI could not be recorded due to the masking of the electrical stimulus artefact, waves eII and eIV were observed in some recordings. Waves eIII and eV were evoked in all cases at 200 CU. In all electrodes, wave eV latencies were seen to prolong as stimulus intensity decreased. ECAP and EABR thresholds and wave eV latencies obtained in EABR thresholds are given in Table 1.

Based on the values given in Table 1, no significant differences were observed between the EABR and the ECAP thresholds in basal, medial and apical regions (p>0.05). Moreover, the significance of the difference between the ECAP and the EABR thresholds was analyzed in terms of the site of the electrode. EABR thresholds were not found significantly different between basal, medial and apical region electrodes (p>0.05), but ECAP thresholds showed significant differences between apical and basal region electrodes (p=0.002). When the difference between EABR wave eV latencies at the threshold level was analyzed with respect to electrode regions, significant difference was found between basal and apical regions (p=0.03).

Evoked auditory brainstem response results according to electrode regions are given in Table 2. Wave eIII and eV amplitudes

Table 1. ECAP and EABR thresholds and wave eV latencies obtained in EABR thresholds

<table>
<thead>
<tr>
<th>Mean±Standard Deviation</th>
<th>ECAP threshold (CU)</th>
<th>EABR threshold (CU)</th>
<th>Wave eV latency at threshold (msec)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Basal electrode</td>
<td>171.5±11.38</td>
<td>169.69±12.17</td>
<td>4.88±0.4</td>
</tr>
<tr>
<td>Medial electrode</td>
<td>169.69±20.32</td>
<td>165.62±16.41</td>
<td>4.8±0.32</td>
</tr>
<tr>
<td>Apical electrode</td>
<td>160.81±20.03</td>
<td>160±15.49</td>
<td>4.53±0.35</td>
</tr>
</tbody>
</table>

ECAP: electrically compound action potentials; EABR: electrically evoked auditory brainstem response; CU: current unit; msec: millisecond
and latencies obtained at 180 CU and 200 CU, and eIII-eV interwave intervals are shown in this table.

An analysis of wave eV latencies at 200 CU and 180 CU intensity as given in Table 2 showed that wave eV latencies obtained at 180 CU did not significantly differ (p>0.05), but wave eV latencies obtained at 200 CU did significantly differ among the electrode regions. These results are given in Table 3.

Differences between EABR wave latencies and amplitudes at 200 CU and 180 CU are given in Table 4. Accordingly, wave eIII and eV latencies and amplitudes were identified to show significant differences at 200 CU and 180 CU stimulation. Only eIII-eV interwave intervals did not show any differences.

Additionally, significant correlation was identified between the EABR and the ECAP thresholds in each of the stimulation regions. Correlation was observed to be greater for the basal region, and uphill in the medial and apical regions. At the same time, EABR and ECAP levels for the medial and the apical regions were observed to have an uphill and moderate correlation (Table 5).

Discussion

In all cases included in this study, high-amplitude eV waves with significant peaks were obtained in the apical region especially at 200 CU. Wave eV latencies evoked by apical electrodes were significantly shorter than those evoked by basal electrodes. Also, ECAP thresholds evoked by apical electrodes were found to be lowest and significantly different from those evoked by basal electrodes.

In our study, we also found that ECAP and EABR thresholds showed an uphill, strong and significant correlation among apical, medial and basal electrode regions. In a study which they conducted with Nucleus CI users, Brown et al. (13) demonstrated that there were no significant differences between ECAP and EABR thresholds.

Hay-McCutcheon et al. (14) have compared the EABR and EAP (Electrically Evoked Whole-Nerve Action Potential) measurements in 10 postlingual adults with Nucleus CI 24R and Nucleus CI 24M implants, and reported to have found significant difference between the EAP and the EABR thresholds in the Nucleus CI 24M users, but no significant difference in Nucleus CI 24R users. It is probable that stimulant parameters and/or demographic characteristics of sample groups are different in studies that report significant differences between ECAP and EABR thresholds and the results were affected by these differences.

In a study which Hughes and Stille (6) evaluated the psychological and physiological measurements of electrical field interaction in CI users, they reported that significantly higher ECAP thresholds were evoked by basal electrodes compared to those evoked by apical electrodes. This result is consistent with our results.

Firszt et al. (2) reported high levels in EABR thresholds obtained from apical electrodes. In another study, the same group of authors studied lateral to medial electrode placement in terms of EABR and reported that the electrode regions showed differences by EABR thresholds. They found that the thresholds obtained from medial electrodes were lower and wave amplitudes were higher (15). In our study, EABR thresholds in each of the electrode regions were evaluated and no significant differences were found.

In our study, we found significant difference between the basal and apical electrodes in wave latencies at EABR threshold level. Significantly earlier wave eV latencies were obtained from apical electrodes than basal electrodes.

In our study, we found significant difference between the basal and apical electrodes in wave latencies at EABR threshold level. Significantly earlier wave eV latencies were obtained from apical electrodes than basal electrodes. Similar results are also found in the literature (6, 15-17). A study investigating the effects of anatomy on EABR reported that wave eIII and eV latencies were significantly affected by the stimulated region, and mean latency levels of wave eV showed an increase of 0.43 msec from the apex to the basal in all electrodes (18). This was explained by the larger diameter of the apical nerve fibers (5). In our study,

### Table 2. EABR latency and amplitude levels obtained at 180 and 200 CU by electrode stimulation regions

<table>
<thead>
<tr>
<th>Mean Standard Deviation</th>
<th>Wave eIII latency (msec)</th>
<th>Wave eIII amplitude (µV)</th>
<th>Wave eIII latency (msec)</th>
<th>Wave eV amplitude (µV)</th>
<th>eIII to eV interwave interval (msec)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Basal electrode</td>
<td></td>
<td></td>
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</tr>
<tr>
<td>200 CU</td>
<td>2.16 ± 0.13</td>
<td>0.21±0.17</td>
<td>4.27±0.3</td>
<td>0.29±0.16</td>
<td>2.15±0.55</td>
</tr>
<tr>
<td>180 CU</td>
<td>2.16 ± 0.09</td>
<td>0.13±0.08</td>
<td>4.35±0.37</td>
<td>0.22±0.11</td>
<td>2.08±0.24</td>
</tr>
<tr>
<td>Medial electrode</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>200 CU</td>
<td>2.07±0.24</td>
<td>0.35±0.13</td>
<td>4.06±0.29</td>
<td>0.43±0.17</td>
<td>2±0.11</td>
</tr>
<tr>
<td>180 CU</td>
<td>2.09±0.14</td>
<td>0.18±0.09</td>
<td>4.22±0.32</td>
<td>0.23±0.08</td>
<td>2.05±0.17</td>
</tr>
<tr>
<td>Apical electrode</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>200 CU</td>
<td>2.04 ± 0.21</td>
<td>0.38 ± 0.19</td>
<td>3.93 ± 0.25</td>
<td>0.4 ± 0.15</td>
<td>1.9±0.15</td>
</tr>
<tr>
<td>180 CU</td>
<td>2.17 ± 0.34</td>
<td>0.24±0.16</td>
<td>4.11±0.17</td>
<td>0.29±0.12</td>
<td>1.93±0.33</td>
</tr>
</tbody>
</table>

CU: current unit; msec: millisecond; µV: microvolt; EABR: electrically evoked auditory brainstem response
wave eV latency at 200 CU showed a significant difference among all stimulation regions. No significant differences were identified among wave eV latencies at 180 CU in terms of electrode regions. There are studies that support (2, 15, 19, 20) this result and advocate the opposite (9, 17, 21, 22) in the literature.

In our study, wave eIII latencies were examined at 200 CU and 180 CU intensities. Wave eIII latency at 200 CU was significantly prolonged in the basal electrode. The difference between the apical and the medial electrodes was not significant. No significant differences were identified among wave eIII latencies at 180 CU in terms of electrode regions. This result may suggest that the stimulation level may have affected wave eIII latency since stimulation at higher intensities can both cause latency and stimulate more central areas. This finding has been previously reported in the literature (23). Guirauda et al. (23) reported that wave eIII latency was significantly affected by the stimulated region, and mean latency levels showed an increase of 0.49 msec from the apex to the basal in all electrodes.

No significant differences were found in our study among apical, medial and basal electrodes in the eIII–eV interval at 200 CU and 180 CU intensities. This result was found consistent with the literature. In a study examining the characteristics of hearing loss and the anatomical effects of auditory pathways with EABR, the effect of the stimulated region on wave eIII–eV interval was not found statistically significant (18). In the light of the data obtained in our study and from the literature, we can say that wave eIII latency is prolonged or shortened parallel to the eV wave latency and this applies to all electrode regions.

In the literature, eII wave latency is reported as 1.30 msec, eIII wave latency as 2.10 msec, and eV wave latency as 3.75 msec at a stimulus level close to the maximum behavioral dynamic range.

Similarly, interwave latencies were reported as 0.80 msec for the eII–eIII interval, 1.60 msec for the eIII–eV interval, and 2.40 msec for the eII–eV interval (2, 16, 24, 25). Wave eV amplitude was defined in the range of 1.00 to 1.46 µV at high stimulus intensity depending on the recording electrode in the implant electrode array (26). In our study, the results closest to these levels were obtained from apical stimulation at 200 CU. Wave eV amplitude at 200 CU did not show a significant difference in the apical and the medial electrodes but decreased significantly in the basal electrode. In our study we also examined wave amplitudes at 180 CU and found no significant differences in wave V amplitudes with respect to electrode regions. Stimulus intensity, as well as electrode placement alter EABR amplitude.

Each of the EABR response parameters—latency, amplitude and morphology—varies as a function of the electrode position. Wave eV latency of the basal region electrode is longer than that of the apical region electrode (27, 28). This result was also found in our study.

In their study they conducted in 2006 with seven adult and seven pediatric Nucleus 24 CI users, Çiprut and Akdağ (29) reported EABR results in which wave eV amplitudes decreased as stimulus levels decreased, and wave morphology was better in apical channels.

That, in our study, a significant peak was observed in wave eV in all cases at 200 CU in the apical regions and a decrease was seen towards the basal in the number of cases at the same intensity, is consistent with the studies reporting that the population of live spiral ganglion cells affect the amplitude and morphology variables in EABR waves (18). When the stimulus level decreased to 180 CU, the number of cases with significant peaks also decreased. The flattened and ovaliform state which wave eV has assumed from the apical to the basal at 200 CU was also observed at 180 CU.

**Conclusion**

In EABR, it will be better to select multiple electrodes from each of the basal, medial and apical regions to record responses specific to each stimulation region. In EABR, the most distinct response is evoked at 200 CU. EABR recording can be started at this intensity level. At the same time, 200 CU was a sufficient level for demonstrating the characteristics of wave latency and wave morphology. EABR or ECAP recording techniques are not suitable substitutes in CI assessment. While ECAP can be preferred for its clinical practicality, EABR may be more suitable for use in clinical trials because of the more qualitative information it provides.
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