Electrophysiological Monitoring During Acoustic Neuroma and Other Posterior Fossa Surgery


ABSTRACT: Techniques used to monitor the function of the seventh and eighth cranial nerves during acoustic neuroma and other posterior fossa surgery are reviewed. The auditory brainstem response (ABR), electrocochleogram (ECochG) and direct recording from the auditory nerve (CNAP) were compared. The best technique is the ECochG, although in many cases, the CNAP should be used as a back-up technique. The CNAP is especially useful for the identification of the auditory nerve. Both can provide real-time feedback on the physiological integrity of the auditory nerve. The ABR may be helpful in monitoring brainstem function. For some procedures, optimal monitoring requires the combined recording of all three techniques.

Monopolar constant-voltage intracranial stimulation of the facial nerve is helpful for the identification and preservation of the facial nerve. Audio monitoring of spontaneous electromyographic activity provides real-time feedback on the effect of surgical manipulation of the nerve. Monitoring of ephaptic transmission in the facial nerve during microvascular decompression for hemifacial spasm aids in the identification of the offending vessel.

One of the principal objectives of posterior fossa surgery is to preserve the function of the seventh (facial) and eighth (auditory) cranial nerves and the brainstem. Intraoperative monitoring can provide real-time feedback to the surgeon on the physiological integrity of these structures, and can reduce the risk of iatrogenic injury. The auditory brainstem response (ABR) was the first technique used to monitor the physiological integrity of the auditory nerve during acoustic neuroma surgery. This technique is non-invasive and easy to use. The early peaks of the ABR are relatively insensitive to the effect of anesthetics. However, the first peak of the ABR may be difficult to identify in the recorded waveform, especially in patients with hearing loss, and therefore, it has been recommended that a more direct method of monitoring peripheral auditory nerve function be utilized. This is of particular importance for intraoperative monitoring. Moller and Jannetta developed a method for recording the compound action potentials directly from the cochlear nerve (CNAP), which provides a large, easy to elicit response. The disadvantage with this technique is that the presence of the electrode in the operative field may be cumbersome. The electrocochleogram (ECochG) which is recorded from an electrode placed in the middle ear avoids this problem. We have examined the relative merits of these three recording techniques (ABR, CNAP and ECochG) for the intraoperative monitoring of the function of the seventh cranial nerve during acoustic neuroma surgery.
recording of auditory and brainstem function during posterior fossa surgery.

During surgery for lesions of the cerebellopontine angle, there is also a risk of injury to the facial nerve. Electrophysiological monitoring has increased the ability of the surgeon to preserve the facial nerve. Rand and Kurze developed a technique for continuous monitoring of mechanical stimulation of the facial nerve by attaching accelerometers developed a technique for continuous monitoring of mechanical monitoring has increased the ability of the surgeon to preserve the facial nerve. Furthermore, the stimulating current might spread to either the spinal accessory or trigeminal nerves, and give false-negative results. For this reason, Delgado and his colleagues introduced intraoperative monitoring of the electromyogram (EMG). Intraoperative photographs were taken of the EMG response from the oscilloscope to determine if there was a change in the latency or amplitude of the response. Although this method permitted identification and mapping of the facial nerve, the stimulus artifact could obscure the waveforms. To overcome this, Sugita and Kobayashi developed a technique for continuous monitoring of mechanical movement of the facial nerve by attaching accelerometers to the orbicularis oculi and oris muscles. Facial muscle movement was then converted to sound through an amplifier and speaker connected to the accelerometer. This system permitted easy identification of the facial nerve and, more importantly, also enabled the surgeon to monitor mechanical movement of the nerve, therefore providing real-time feedback indicating mechanical injury to the nerve during dissection. Moller and Jannetta provided the next refinement by developing a system combining sound monitoring of spontaneous EMG activity, and monitoring of EMG activity in response to direct intracranial stimulation of the facial nerve. This was accomplished by introducing a gating circuit to remove the stimulus artifact thereby allowing facial muscle action potentials to be monitored with a loudspeaker. The present report also contains our experience with the intraoperative monitoring of facial nerve function during posterior fossa surgery.

Electrophysiological monitoring of the facial nerve has also become an important intraoperative tool for microvascular decompression of the facial nerve for hemifacial spasm. Once the facial nerve is decompressed in these patients, the ephaptic transmission disappears, and therefore this method permits the easy identification of the offending vessel, and improves the effectiveness of the procedure. This report also contains a description of our experience in monitoring facial nerve function during microvascular decompression for hemifacial spasm.

### MATERIALS AND METHODS

#### Clinical Material

Electrophysiological monitoring was performed on 19 patients undergoing microsurgery for the removal of acoustic neuromas or other cerebellopontine angle (CPA) tumors, or for microvascular decompression of the VII cranial nerve for hemifacial spasm (Table 1). The management of these patients, including the surgical approaches used for the removal of acoustic neuromas and other CPA tumors, have been described elsewhere. Microvascular decompression for hemifacial spasm was performed using a posterior fossa approach.

#### Anesthesia

Electrophysiological monitoring of auditory function during posterior fossa surgery has few anesthetic limitations. Body temperature must be monitored because hypothermia will cause an increase in the latency of the ABR responses. Barbiturates, inhalation anesthetics, and muscle relaxants have a small but definite effect upon the ABR. The early peaks of the ABR are much less affected than the later peaks. In our unit, anesthesia is maintained with isoflurane and nitrous oxide, and fentanyl is administered as required. Manninen et al examined the effect of isoflurane on the ABR and found that all the ABR peaks were affected, although the more central the generator sites of these

### Table 1

<table>
<thead>
<tr>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>APF</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>No</td>
<td>No</td>
</tr>
<tr>
<td>2</td>
<td>ATA</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>No</td>
<td>Yes</td>
</tr>
<tr>
<td>3</td>
<td>ATA</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>No</td>
<td>No</td>
</tr>
<tr>
<td>4</td>
<td>CPA</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>No</td>
<td>No</td>
</tr>
<tr>
<td>5</td>
<td>CPA</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>No</td>
<td>Yes</td>
</tr>
<tr>
<td>6</td>
<td>APA</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>Yes</td>
<td>Yes</td>
</tr>
<tr>
<td>7</td>
<td>APA</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>Yes</td>
<td>Yes</td>
</tr>
<tr>
<td>8</td>
<td>APA</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>Yes</td>
<td>Yes</td>
</tr>
<tr>
<td>9</td>
<td>APA</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>Yes</td>
<td>Yes</td>
</tr>
<tr>
<td>10</td>
<td>CPA</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>Yes</td>
<td>Yes</td>
</tr>
<tr>
<td>11</td>
<td>APA</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>Yes</td>
<td>Yes</td>
</tr>
<tr>
<td>12</td>
<td>APA</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>Yes</td>
<td>Yes</td>
</tr>
<tr>
<td>13</td>
<td>APA</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>Yes</td>
<td>Yes</td>
</tr>
<tr>
<td>14</td>
<td>APA</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>No</td>
<td>Yes</td>
</tr>
<tr>
<td>15</td>
<td>APA</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>Yes</td>
<td>Yes</td>
</tr>
<tr>
<td>16</td>
<td>APA</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>Yes</td>
<td>Yes</td>
</tr>
<tr>
<td>17</td>
<td>APA</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>No</td>
<td>Yes</td>
</tr>
<tr>
<td>18</td>
<td>APA</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>Yes</td>
<td>Yes</td>
</tr>
<tr>
<td>19</td>
<td>APA</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>*</td>
<td>Yes</td>
<td>Yes</td>
</tr>
</tbody>
</table>

**ATA** Translabyrinthine approach for the removal of large acoustic neuromas (greater than 2 cm)

**APF** Posterior fossa approach for the removal of small acoustic neuromas (less than 2 cm)

**CPA** Other cerebellopontine angle tumours

**FD** Facial nerve decompression

* Test performed
components, the greater the effect. Our regimen avoids the administration of long-lasting muscle relaxants and thus allows the facial EMG to be recorded. Short-acting muscle relaxants (e.g., Atracurium) are used for induction, the effects of which have disappeared by the time the recording of the EMG activity is required. After the tumor has been removed, or the facial nerve decompressed, longer acting muscle relaxants may be given.

Monitoring of Auditory Functions

The Cadwell 8400 system was used for monitoring auditory function as described in (a), (b) and (c) below.

(a) Auditory Brainstem Response

The ABR was recorded preoperatively, intraoperatively and postoperatively. Preoperatively and postoperatively, 70 dB nHL rarefaction clicks (100 usec) were presented through a moulded ear insert at a repetition rate of 11.1/s and 44.4/s. A plastic tube, 1.9 mm in diameter and approximately 32 cm in length, was used to conduct sound from the speaker to the earmould. Response threshold was determined by presenting 50, 30, and 10 dB nHL clicks at a repetition rate of 44.4/s. Each trial consisted of recording the averaged response to 2000 stimuli presented monaurally, and then the trial was replicated. Masking noise was presented to the ear contralateral to the stimulus at an intensity 40 dB less than the intensity of the click stimulus. Gold-plated cup electrodes were attached to the scalp with saline gel and collodion-impregnated gauze, and placed at Cz, M1, M2, the Inion and 2 cm above each auricle (A). Four channels of activity were recorded with the following derivation: Cz-M1, Cz-M2, Cz-Inion, Cz-Ai (i = ipsilateral). A ground electrode was taped to the shoulder. The interelectrode impedences were less than 5 KOhms.

Negativity at the vertex was plotted in the upward direction. The EEG was amplified with a bandpass of 100 to 3000 Hz. The sweep time was altered to 20 ms. If there was no reproducible response, the sweep time was altered to 20 ms. Intraoperatively, 70 dB nHL stimuli were presented at a rate of 44.4/s, but the other stimulus and recording parameters were identical to those used preoperatively.

(b) Direct Recording of Compound Nerve Action Potentials from the Auditory Nerves

The procedure used to record the CNAP was similar to the technique developed by Moller and Jannetta. A cotton wick electrode was constructed by removing the end 6 mm of insulation from Ag5T wire, to which cotton wool was sutured to the bent, exposed wire. The wire was then attached to insulated copper wire which was connected to the non-inverting input of the preamplifier. The reference electrode was constructed with a metal 25 gauge hypodermic needle connected to insulated copper wire, which was connected to the inverting input of the preamplifier. The needle electrode was inserted into muscle at the operative site, and the cotton-wick electrode was sutured to a Sugita retractor and held in place at the operative site.

Auditory click stimuli were presented through earmoulds, and masking noise was presented to the contralateral ear at an intensity 40 dB less than the stimulus used to elicit a response. For monitoring purposes, 40 click stimuli were presented (70 dB nHL) at a presentation rate of 22.2/s. The system bandpass was 10-3000 Hz and the sweep time was 10 ms. A-1 ms sweep delay was used occasionally to monitor stimulus artifact. Figure 1 illustrates our method of monitoring auditory function with this technique.

(c) Electrocochleogram (ECochG)

An electrode identical to that used to record the CNAP from the auditory nerve was used to record the ECochG except that Pt5T wire was used because it is more malleable than Ag5T and is easier to anchor in the external auditory meatus. A myringotomy was performed, through which the electrode was placed against the promontory, and the electrode was held in position by the moulded ear insert (Figure 2). The impedance of the transtympanic electrode was 40 to 50 KOhms. A needle EEG electrode placed on the face was used as the reference electrode. The effects of varying the stimulus intensity, polarity and presentation rate were examined, and the number of averages required to obtain a satisfactory signal-to-noise ratio, and optimal filter settings were determined.

Monitoring of Facial Nerve Function

(a) Facial Monitoring During Acoustic Neuroma Surgery

After the patient’s head was secured in the head holder, a hand-held stimulator with metal probes was placed on the skin overlying the marginal mandibular branch of the facial nerve. The nerve was stimulated, a muscle twitch observed in the orbicularis oris muscle, and then two platinum needle electrodes were placed in the muscle. A second set of electrodes were placed in the orbicularis oculi muscle after localization by stimulating the zygomatic branch of the facial nerve. The electrode wires were taped to the face. Constant-current stimuli were used to localize the muscle twitch with the Cadwell-8400 signal averaging system being used for these preliminary procedures. Thereafter, the Grass-Moller NL-1 facial nerve locator and monitor were used for monitoring as described by Moller and Jannetta. One electrode from each muscle was inserted into the input probe of the Grass locator. The remaining electrodes were used as back-up electrodes. A ground
electrode was placed in the forehead. The responses were differentially amplified and displayed on an oscilloscope. The system bandpass was 0.3-3000 Hz. The responses were also audio-amplified, providing immediate feedback about mechanical stimulation of the nerve. For identification purposes, monopolar constant-voltage stimulation was applied through a hand-held nerve locator unit (Figure 3). A maximum stimulus intensity of .7 V was applied intracranially. Stimuli of 200 usec duration were presented at a rate of 10/s.

(b) Facial Nerve Monitoring During Microvascular Decompression of the Facial Nerve for Hemifacial Spasm

Needle stimulating electrodes were placed over the zygomatic and marginal mandibular branches of the facial nerve. Electromyographic activity was recorded with needle electrodes placed in orbicularis oculi and mentalis muscles. Constant-voltage stimuli (250 usec duration) were presented at a rate of 11.1/s. The system bandpass was 1-3000 Hz.

RESULTS

(a) Auditory Nerve Monitoring

Monitoring of the ECoG and CNAP has been of great value intraoperatively in patients with acoustic neuromas because both provide real-time feedback to the surgeon on the physiological integrity of the eighth cranial nerve. Intraoperatively, the ABR has been less useful as a technique to provide immediate feedback. However, the ABR can be used as a back-up system. Furthermore, if the lesion is close to the brainstem, CNAP recordings are difficult to obtain and the ABR may then be used for monitoring. The ABR can also be used as a measure of brainstem function. For example, during the removal of a...
large acoustic neuroma, one patient had extensive blood loss which caused a marked decrease in amplitude and increase in latency of wave V of the ABR recorded in response to auditory stimuli delivered to the ear contralateral to the surgery. This was a helpful warning of brainstem dysfunction.

Direct recording of the CNAP from the auditory nerve was found to be very useful in many cases of acoustic neuroma in which hearing preservation was attempted. Figure 4 is from a patient with a large acoustic neuroma in whom the CNAP gradually increased in latency and decreased in amplitude, and then was lost completely. Post-operatively, hearing was absent on the operated side. One of the problems with this technique is the presence of the electrode in the operative field. The primary purpose of this method of monitoring auditory function is to identify the cochlear nerve. With this technique, the surgeon can differentiate tumor from nerve. The CNAP recording can also be used as a back-up method to other methods of continuous monitoring of auditory function.

The ECochG proved to be the best method of monitoring auditory nerve function. The cotton-wick electrode was easy to use and out of the operative field, and the recording electrode was secure when anchored in place with the ear-insert. To date, there have been no problems with the loss of the ECochG recording electrode due to displacement or high electrode impedance. However, because of these possibilities, back-up systems of monitoring such as the CNAP should be used.

The effect of stimulus polarity on the ECochG was also examined intraoperatively, and Figure 5 illustrates the importance of this relationship. Occasionally it was difficult to identify the N1 component. Because the phase of the CM is opposite if condensation or rarefaction clicks are presented, the CM can be removed by presenting alternating stimuli. The resulting waveform clearly demonstrated the N1 component which represents activity generated from the auditory nerve.

The effect of stimulus intensity on the ECochG was also examined intraoperatively. When alternating clicks are presented, the cochlear microphonic (CM) is subtracted from the waveform allowing examination of the N1 component. As the stimulus intensity was increased, the amplitude of the N1 component increased and the latency decreased.

The effect of stimulus presentation rate on the ECochG is shown in Figure 6. The response can be recorded at high presentation rates (88.8/s), but as the rate of stimulus presentation was increased, the amplitude of the N1 component decreased. However, even at fast presentation rates, a clear N1 component was evident. The CM response was also present at high presentation rates. The CM response was a following response. The number of cycles observed in the CM increased as the presentation rate increased. Because the ECochG could be recorded with a small number of averages (2-10) at a high presentation rate, real-time monitoring of the function of the auditory nerve was obtained.

The effect of altering the high-pass and low-pass filter settings was examined intraoperatively. We found that the high-cut filter setting should not be lower than 3000 Hz, and the low-cut filter setting should be 10 Hz.

During the removal of the posterior wall of the internal auditory canal with the drill in the suboccipital approach to acoustic neuromas, it is too time consuming to pause long enough to record the ABR because of the number of responses which have to be averaged. In contrast, the ECochG can be recorded

![Figure 4](https://www.cambridge.org/core/core-image.png)  
**Figure 4** — Intraoperative monitoring of the compound nerve action potentials (CNAP) recorded from the auditory nerve. Hearing preservation was attempted during the removal of a large acoustic neuroma. Shown on the right of each waveform is the time of recording. At 11:40 a reproducible response was evident, but at 12:40 the amplitude of the response decreased and the latency increased. This trend continued and at 3:35 there was no response present.

![Figure 5](https://www.cambridge.org/core/core-image.png)  
**Figure 5** — The effect of stimulus polarity on the ECochG examined intraoperatively. Each waveform is the averaged response to 40 stimuli presented at a rate of 22.21/s at an intensity of 70 dB nHL. The tracing at the top of the figure was recorded in response to condensation clicks; the middle tracing was recorded in response to rarefaction clicks; and the lower tracing was recorded in response to alternating clicks. Note the phase reversal of the CM when condensation clicks were presented as compared to rarefaction clicks. Note the removal of the CM component when alternating clicks were presented.
Figure 6 — The effect of stimulus presentation rate on the ECochG recorded intraoperatively. Each represented waveform was elicited by presenting 40 rarefaction clicks at an intensity of 70 dB nHL. The rate of stimulus presentation is listed on the right side of the figure.

in less than a second, which allows recording of the response between drilled sessions. It should be noted that waveforms should be interpreted cautiously at this time because an increase in latency and decrease in amplitude of the ECochG has been observed. This change may be caused by a temporary shift in the response threshold.

Figure 7 shows the ECochG and the CNAP from a patient with bilateral acoustic neuromas in whom hearing was preserved during the removal of a small acoustic neuroma less than 1 cm in diameter. During the dissection an anomalous artery entering the temporal bone was retracted and caused an abrupt loss of both the CNAP and ECochG responses. The surgeon was informed and the operation was halted. Ten minutes later, the response returned. The method of tumour removal was altered and the patient awoke with some preservation of hearing. Because of the complete yet reversible loss of the ECochG, this suggests reversible cochlear ischemia.

(b) Facial Nerve Monitoring

The audio monitoring of mechanical stimulation of the facial nerve provided real-time feedback to the surgeon and was very useful during intracranial dissection of the facial nerve, especially in large acoustic neuromas. The differentially amplified EMG activity was large in amplitude and easy to record. The placement of multiple back-up electrodes in the facial muscles was helpful because the needle electrodes can dislodge. A portable impedance meter was helpful to test contiguity. In the development of our monitoring methods, a variety of hand-held facial nerve stimulators were constructed and tested. Both monopolar and bipolar constant-current and constant-voltage methods of stimulation for identifying the nerve were evaluated. The commercially available Moller stimulator from Grass Instruments proved to be the best for two reasons: the probe tip was malleable, and allowed enhanced adaptability in the operative field; and the monopolar constant-voltage stimulation provided consistent results.

Electrophysiological monitoring during microvascular decompression for hemifacial spasm enhanced the accuracy of the procedure (Figure 8). The disappearance of the ephaptic responses was a helpful indicator of the adequacy of decompression.

Figure 7 — The top three tracings illustrate the ECochG recording and the lower three tracings illustrate the CNAP recordings. The tracings on the left side of the figure are the baseline recordings. The tracings in the centre were recorded during the manipulation of an anomalous artery to the temporal bone when both the ECochG and CNAP readings were lost. The tracings on the right were recorded after the surgical team halted the operation for ten minutes after which the response returned.
Figure 8 — Electrophysiological monitoring during microvascular decompression for hemifacial spasm. The top tracings illustrate EMG activity recorded from the mentalis muscle. The lower tracings illustrate EMG activity recorded from orbicularis oculi. Both sets of tracings were recorded in response to stimulation of the marginal mandibular branch of the facial nerve. Before microvascular decompression, a direct response was recorded from the mentalis muscle and an indirect response was recorded from orbicularis oculi. After the decompression there was no longer an indirect response recordable from the orbicularis oculi muscle.

DISCUSSION

The ABR is an accurate diagnostic test in conditions such as acoustic neuroma where it is abnormal in at least 90% of cases. However, for a number of reasons, its value in the operating room is limited. Firstly, to acquire an adequate signal-to-noise ratio requires the averaging of thousands of responses, which is time consuming. Secondly, the later components of the ABR are affected by hypothermia and some anesthetics. Finally, during intraoperative monitoring of the ABR, wave V has been used to identify the response, but intraoperative changes in wave V do not correlate well with post-operative hearing preservation. Accordingly, attention has been directed towards monitoring the earlier responses of the auditory system, especially at the level of the auditory nerve because these earlier signals are more resilient to changes in body temperature and are less affected by anesthetic level.

There is a wealth of literature on the clinical use of the ECochG. The ECochG was first recorded intraoperatively in the 1960's during middle ear surgery. Because of the work of Levine and his colleagues there has been a reawakening of interest in the intraoperative recording of the ECochG. The stimulus/intensity function of the N1 component of the ECochG which we have observed intraoperatively is similar to that described by Eggermont. The amplitude of the N1 component decreases and the latency increases as the stimulus intensity is increased. At the highest intensities, corresponding to the H area of the amplitude/intensity function, the N1 component may have three peaks. At lower intensities the second peak is the most prominent. As the stimulus rate is increased, the N1 response amplitude decreases and the latency increases. However, the N1 component is still present at fast rates (200/S). When the rate is greater than 80/s, the summing potential may contaminate the recordings. Therefore, we recommend a rate of 44/s. Until the clinical usefulness of the CM is determined, alternating click stimuli should be applied to remove the CM, which can, as we have demonstrated, contaminate the N1 component of the waveform.

The ECochG has several advantages, including ease of recording and the absence of the electrode in the operative field. It has also been reported that the ECochG is not altered by anesthetic levels. However, there are several aspects of the intraoperative recording of the ECochG which require further research. Firstly, the method of recording might be improved. For example, Stone et al reported that the ECochG can be recorded from a low impedance electrode placed near the opening of the external auditory canal, rather than transtympanically. Although transtympanic recordings are larger, they require penetration of the tympanic membrane. Further research is required to determine the effectiveness of the external electrode technique in the operating room. A second area requiring further research is the CM response, which may provide an accurate assessment of the inner hair cells of the cochlea. To date, most of the work has focused on the N1 component. Because of the periodicity of the CM component, frequency based techniques such as Fourier analysis may be used to analyze this response. Indeed, response detection methods may be significantly improved by using optimal digital filtering and higher level statistical techniques, such as the Hotellings t2 test or "phase coherence"." Finally, the effects of anesthetics and temperature on the ECochG should be more rigorously examined.
Use of the ECochG alone has several limitations. Firstly, the recording electrode may be dislodged away from the promontory during the operation, and would be difficult to replace. Thus, whenever possible, the CNAP and the ABR should also be recorded as back-up methods. Another potential problem with the ECochG is that the N1 component is thought to be generated from the distal portion of the auditory nerve and accordingly, if the tumour is proximal to the generator site of the N1 component, the ECochG may not detect an injury to the nerve near the tumour. This problem may also be minimized by the additional recording of the CNAP and ABR.

We conclude that the peripheral auditory nervous system can be accurately monitored with the combination of the ECochG and the CNAP. With small acoustic neuromas, combined recording provides excellent monitoring for operations designed for hearing preservation (Table 1). For monitoring during the removal of large acoustic neuromas, a different approach is required because hearing preservation is no longer the primary objective of auditory evoked potential monitoring. With large tumours, the function of the facial nerve and the brainstem are in jeopardy and electrophysiological methods of monitoring these structures are required. The ABR can be used for monitoring brainstem function, although the ABR is not sensitive to damage to areas of the brainstem other than the ascending auditory pathway. Indeed, Piatt et al. recommended that the sensitivity of electrophysiological monitoring would be enhanced if somatosensory evoked potentials were recorded in conjunction with the ABR. Further work is required in this important area.

New surgical approaches for the removal of large acoustic neuromas have improved the likelihood of preservation of the facial nerve, and the same is true for electrophysiological monitoring. Electrical stimulation helps to identify the facial nerve, although there is controversy about the optimal method for intracranial stimulation of the facial nerve. Babin et al. recommended a bipolar constant-current stimulator for localization of the facial nerve, whereas Moller et al. preferred monopolar constant-voltage stimulation. Prass described stimulation with flush-tip probes and found no advantage to either constant-current or constant-voltage stimulation. We found that the monopolar constant-voltage technique developed by Moller is reproducible, easy to use, and very helpful for preserving facial nerve function (Table 1). Moller and Jannetta also recommended this technique for stimulating tumour tissue prior to removal if there is doubt about the location of the facial nerve. Finally, we also recommend audio monitoring of spontaneous EMG activity because it can provide real-time feedback on the effect of surgical manipulation on the facial nerve.

Idiopathic hemifacial spasm has been attributed to compression of the root entry zone of the facial nerve by a blood vessel. Intraoperatively, it is possible to determine the offending vessel electrophysiologically. The exact mechanism by which impulses are transmitted along non-stimulated branches of the facial nerve when another branch is stimulated is unknown. Moller and Jannetta reviewed this phenomena. Transmission is either ephaptic or through the facial motor nucleus. Whenever the abnormal activity is generated it is relieved after decompression. Although this abnormal transmission is mainly caused by compression due to blood vessels, in certain cases, other lesions of the cerebellopontine angle may be the cause of the compression. This mode of monitoring may help to differentiate if the lesion is vascular. Because there is a risk of injury to the auditory nerve during microvascular decompression for hemifacial spasm, auditory function should also be monitored during these procedures.

**CONCLUSION**

Electrophysiological monitoring in the operating room is a rapidly evolving field. For each sensory modality, methods of monitoring are being developed and improved on. These techniques can enhance the safety and accuracy of many procedures. Monitoring of auditory nerve and facial nerve functions exemplifies the ability of these tools to improve both safety and accuracy during acoustic neuroma and other posterior fossa surgery.

**ACKNOWLEDGEMENTS**

The authors would like to express their thanks to the following: Ms. L. Kennedy for preoperative and postoperative audiological testing; Dr. R. Harrison for providing helpful comments on the methodology for recording the electrocochleogram; Mr. J. Loukides for building the cotton-wick electrodes; and Miss C. Dickson for secretarial assistance.

**REFERENCES**