SHORT REPORT

Neurological complications of anterior spinal surgery for kyphosis with normal somatosensory evoked potentials (SEPs)

L Pelosi, A Jardine, J K Webb

Abstract

We report a case of neurological complications of anterior release for correction of kyphosis. After the operation, the patient had pyramidal weakness and decreased pain sensation below T5, whereas light touch, proprioception and vibration sensation were intact. Clinical and neurophysiological findings in this patient suggested a partial lesion of the spinal cord probably due to ischaemia in the territory of the anterior spinal artery. Intraoperative and postoperative tibial nerve SEPs remained normal, which stresses the need for recording from the motor pathways.

Keywords: spinal surgery; SEPs; motor deficits

We report a case of a postoperative partial spinal cord syndrome, affecting predominantly pyramidal and spino-thalamic pathways below T5, which complicated an anterior spinal release for kyphosis and was associated with normal SEPs.

Case report

A 21 year old male patient was scheduled for a two-stage operation consisting of a thoracoscopic release followed by a posterior fusion for correction of Scheuermann's kyphosis. He had been referred with a 5 year history of progressive thoraco-lumbar back pain which had recently began to radiate down the posterior aspect of both legs associated with intermittent paraesthesiae and numbness in his legs. Preoperative neurological examination was normal, but reflexes were brisk both in the upper and lower limbs.

The anterior release was carried out via a right thoracoscopic approach with SEP monitoring. Segmental vessels were ligated from T6 to T11. Preoperative and postoperative tibial nerve SEPs remained normal, which stresses the need for recording from the motor pathways.

Following 150 µg fentanyl, anaesthesia was induced with 200 mg of propofol. Intubation was performed after paralysis with 10 mg of vecuronium and anaesthesia maintained with 50% nitrous oxide/oxygen and isoflurane at 1.2% end tidal with supplemental narcotic and relaxant as required. Mechanical ventilation was adjusted to maintain end tidal carbon dioxide at 4.8% and mean arterial pressure (MAP) held between 75 and 80 mm Hg (80% preoperative).

During surgery the thoracoscopic sites remained ischaemic and the spinal cord was exposed. The anterior release was carried out via a right thoracoscopic approach with SEP monitoring. Segmental vessels were ligated from T6 to T11. Preoperative and postoperative tibial nerve SEPs remained normal, which stresses the need for recording from the motor pathways.

Somatosensory evoked potentials (SEPs)

Preoperative and postoperative SEPs to stimulation of the tibial nerves at the ankle (square-wave electrical pulses of 0.1 msec duration, 3Hz, intensity adjusted to produce a visible plantar flexion of the foot) were recorded via a pair of silver discs at Cz and Fz (10-20 International System). Bandpass filters were 30-5000 Hz and analysis time 100 ms.
Intraoperative SEPs to alternate stimulation of the tibial nerves at the ankle (7Hz, 0.1 ms duration, intensity adjusted as above prior to administration of muscle relaxants) were derived from a cervical-to-scalp montage (the cathode over the C2 and C5 spinous processes and the anode at Fz) using silver disc electrodes. Surface recordings of the peripheral nerve responses at the knee were obtained simultaneously to ensure adequacy of stimulation. SEPs were recorded continuously during the first two hours of the operation, following which monitoring was discontinued as no instrumentation and corrective manoeuvres potentially hazardous for the spinal cord were to be implemented. SEP monitoring was resumed at the end of the operation before closure.

Intraoperative SEPs remained normal and symmetric throughout the monitoring time. SEPs recorded at the end of the operation were normal with no significant change of amplitude and latency relative to the baseline responses (fig A). Both the preoperative SEPs and the SEPs recorded the day after the operation when the patient was severely disabled were normal (fig B).

Discussion
In our patient, thoracoscopic release for correction of kyphosis was complicated by a partial spinal cord lesion affecting pyramidal and spino-thalamic pathways below T5, but sparing posterior sensory cord functions. We could not document the time when the spinal cord damage occurred, because all vital parameters remained unchanged throughout the operation, except for a sudden decrease of the CFAM amplitude. We could identify no reason for this change, but it is possible that damage to the spinal cord might have begun at that time. Immediate postoperative examination was apparently normal, suggesting that the pathological process evolved over a period of time. Again, no change of vital parameters occurred at the time when the neurological deficits became manifest. Three delayed neurological complications following spinal surgery, two of which purely motor, have been mentioned but not further discussed, in a previous paper. Delayed postoperative paraplegia has also been reported in a 46 year old woman following revision of scoliosis surgery. Neurological deficits begun three and a half hours after skin closure and coincided with a fall in blood pressure probably caused by hypovolaemia. Interestingly, in the same case report, intraoperative falls of blood pressure to the same value blamed postoperatively for cord malfunction were not associated with SEP change.

(A) Sequential recordings of intraoperative SEPs to alternate stimulation of the left and right tibial nerves at the ankle. Surface recording from C2 (2nd cervical spinous process) - Fz. Note that SEPs recorded at the end of the operation (last two traces on each side) were normal. (B) Postoperative cortical SEPs (Cz-Fz) to stimulation of the tibial nerves (TN). The patient had severe motor deficits at the time when these SEPs were recorded.
The distribution and type of neurological deficit in our patient suggest a vascular mechanism. Ischaemia of the anterior spinal artery at thoracic level would be consistent with a partial lesion affecting the anterolateral and central cord. The posterior third of the cord, which has an independent vascularisation via the posterior spinal arteries, would be expected to be spared. In the thoracic region, the anterior cord is particularly vulnerable to ischaemia, because of the narrowing of the anterior spinal artery and the relative paucity of anastomotic radicular branches. The anterior spinal artery syndrome is a complication of surgery on the thoracic and abdominal aorta and has been reported as a rare complication of anterior spinal surgery.

It is possible that other case reports of predominantly motor postoperative sequelae with preserved intraoperative SEPs were also due to a vascular mechanism. As the sensory pathways which generate the SEPs traverse predominantly the posterior cord, it is not surprising that SEPs, both intraoperative and postoperative, remained normal in our patient. There is the possibility that we have missed a transient change of the SEPs which might have occurred during the period of time when monitoring was discontinued. However, normal SEPs were recorded both at the end of the operation and postoperatively when the patient had severe motor deficits.

Recording from the motor pathways should be expected to offer a better chance of detecting and, hopefully, preventing motor deficits due to anterior spinal artery syndrome. Furthermore, studies in non-human primates have shown that the corticospinal pathways are far more sensitive to ischaemia than the sensory tracts. For moderate degrees of ischaemia, profound attenuation of the motor responses could occur without changes in the SEPs. In the case report by Ginsburg and colleagues, the patient awoke paraplegic from posterior instrumentation despite normal intraoperative SEPs. Changes of the SEPs became manifest hours later. Methods for intraoperative assessment of the spinal motor pathways have been developed and improved in recent years, but various technical difficulties have prevented their widespread use. However, the case we report clearly demonstrates that major motor disability can be associated with entirely normal SEPs. Thus, SEPs cannot be relied upon to detect motor dysfunction.

Neurophysiological monitoring is generally performed during posterior instrumentation in order to prevent damage to the cord from corrective manoeuvres. However, an ischaemic lesion can also occur independently at any stage in both the anterior and posterior procedures. This case stresses the importance of monitoring motor function during anterior procedures.

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