Controversies in the anesthetic management of lumbar drains for endovascular aortic surgery

Sarah Marie Cardillo and Wendy K. Bernstein*
Department of Anesthesiology, University of Maryland School of Medicine, USA

Abstract
Lumbar drain insertion is an integral aspect of perioperative care for patients undergoing thoracoabdominal aortic aneurysm (TAA) repair. It has been shown to reduce incidence of paraplegia when both open and endovascular technique is used. Controversy exists in terms of optimal management of these catheters. This commentary aims to present data as well as experienced recommendation in their use.

Despite the increased use of endovascular techniques for thoracoabdominal aortic aneurysms (TAA), the risk of paraplegia after repair remains a significant concern for both the surgical and anesthesiology teams. Spinal cord ischemia or infarction with subsequent neurologic dysfunction occurs with an incidence of 4-13% after thoracic endovascular aortic repair (TEVAR) [1-3]. Decreased spinal cord perfusion persists postoperatively when episodes of hypotension, hemorrhage and elevated cerebrospinal fluid (CSF) pressure occur, which further increases likelihood of paraplegia after TEVAR [4].

The highest risk factors for paraplegia after TEVAR are the extent of endovascular stent coverage and prior distal aortic repair [5]. Increasing spinal cord perfusion through augmentation of arterial pressure, lumbar CSF drainage and reattachment of segmental arteries has been shown to reduce the incidence of paraplegia. Spinal cord perfusion pressure is a function of lumbar CSF pressure subtracted from mean arterial pressure (MAP). Lumbar CSF drainage effectively improves spinal cord perfusion pressure via reduction of CSF pressure. Two separate metaanalyses including 3 randomized controlled trials have demonstrated the efficacy of controlled drainage of CSF for prevention of neurological injury [6,7].

The placement of lumbar CSF drains is relatively safe and recommended for TEVAR. However, contraindications to placement include pre-existing coagulopathy, elevated intracranial pressure, and frequently, emergent surgery [8]. The procedure itself carries potential side effects, including intracranial hemorrhage (ICH) and neuroaxial hematoma. The American Society of Regional Anesthesia and Pain Medicine (ASRA) advises that instrumentation of the neuraxis be avoided in patients with preexisting coagulopathy, which may increase risk of bleeding with intrathecal catheter insertion. This risk is elevated with recent or concurrent administration of antplatelet agents or anticoagulants. In addition, guidelines state that the time from the procedure and anticoagulation dosing should exceed 60 minutes and that the lowest dose of heparin be administered [9].

CSF drainage can be performed prior to the operation in high risk patients or after the procedure if symptoms of spinal cord ischemia occur. The timing of placement varies by center. Benefits of pre-induction placement include increased length of time prior to anticoagulation as well as knowledge of the patient’s pain or paresthesia during placement, which could indicate potential nerve injury. In patients prescreened as high risk or difficult placement, it may be beneficial to place the lumbar drain well before the start of the case when alternative methods such as fluoroscopy can be utilized. Unfortunately, such management requires the patient to be hospitalized preoperatively which increases cost. Furthermore, prolonged catheter insertion can also place the patient at increased risk for infection [10].

The drain should be placed at the L3-4 or L4-5 level in either the seated or lateral decubitus position. Positioning is at the discretion of the anesthesiologist, as both positions offer advantages. The seated position allows for easier palpation of spinous processes through a midline incision, which avoids the more lateral epidural plexus. In addition, the improved lumbar flexion allows potentially more space for needle insertion. Thus, the risk for bloody tap is decreased. On the other hand, the lateral decubitus position limits hydrostatic pressure of CSF and therefore the potential risk of over-draining CSF. The drain should be inserted 8-10 cm, however depths as great as 30 cm have been reported [11]. Although deeper insertion minimizes the risk of inadvertent removal, it may also contribute to paresthesias and possible nerve root injuries [8].

Should a bloody tap be encountered during placement, the appropriate response remains unclear? Traumatic instrumentation of the neuraxis has been associated with up to 50% of neuroaxial hematomas [8,9]. The occurrence of neuroaxial hematoma is rare. In a case series of 162 patients undergoing full heparinization for extracorporeal circulation, there were no spinal hematomas [12]. Another retrospective study, however, reported two spinal hematomas in sixty five patients undergoing TAA repair [11]. At ours and other institutions, if a bloody tap is encountered, the practice is to discontinue...
drainage and proceed with placement at another level. If bloody tap results a second time, case cancellation is considered after discussion with the operative team [13]. It has been suggested that the case be delayed up to 24 hours, however, based on ASRA guidelines this is only indicated for cases involving full heparinization for cardiopulmonary bypass, and otherwise there is no data supporting case cancellation. Animal models have demonstrated a decreased risk for ischemic injury when goal CSF pressure is below 10 mmHg [6]. Studies finding CSF drainage ineffective limited the total CSF drained to 50 mL as well as allowing the CSF pressure to elevate above 10 mmHg [6]. Continuous monitoring intraoperatively with intermittent drainage allows awareness of sudden waveform disruption and possible drain occlusion. It also avoids high volume or rate drainage and potential intracranial hypotension. Protocols for CSF drainage from the lumbar drain range from 10 to 20 mL/hour depending on patient condition [8].

Management postoperatively requires vigilance in serial neurologic examination. In asymptomatic patients, management goals of CSF pressure should continue to be below 10 mmHg with a rate limit of 10-15 mL/h. Patients with neurologic deficit may require drainage down to 5 mmHg or up to 20 mL/hour, which has been shown to improve symptoms [8,13]. At our institution, the catheter is opened to passive drainage for CSF pressures greater than 10 mmHg, and pressure is then reassessed after each 15 mL of CSF drained. Maximum CSF drainage rate is 20 mL/hour and up to 250 mL/day. If the pressure is persistently elevated after 150 mL are drained, the drain position is evaluated and the transducer re-zeroed. If the patient reveals any neurologic signs or symptoms, the catheter is drained to below 10 mmHg coupled with blood pressure augmentation to a MAP >90 mmHg.

Blood in the draining CSF is another issue that may be encountered. This may be indication of an intracranial bleed, the risk of which is associated with excessive CSF drainage and development of intracranial hypotension [14,15]. Bloody drainage has been shown to be a poor marker of an epidural cord hematoma [10]. Should a change in neurologic status accompany blood within the drain, recommended workup includes immediate imaging [8,13]. At our institution, brain imaging is additionally recommended for asymptomatic persistent bloody CSF drainage lasting more than 4 hours.

Based on current literature, drainage should be discontinued after 2-3 days if there is no sign of spinal cord ischemia. After this period, a fibrotic response will have occurred around the catheter, which seals the puncture site. Clamping the drain prior to removal may lead to greater chance of CSF leak secondary to elevated CSF pressure [13]. However, our institution recommends that the drain be clamped for 24 hours and the patient monitored for neurologic symptoms prior to removal. If a CSF leak occurs, conservative measures such as bed rest, limiting head of bed elevation and hydration are recommended [13].

Lumbar catheters are usually removed if there is no sign of neurologic dysfunction or spinal cord ischemia. Removal of the lumbar drain should always be performed by an experienced practitioner who is familiar with the catheters. Pulling the catheter out slowly and steadily will reduce the risk of catheter fracture. Positioning in the lateral decubitus position with hip and back flexion increases the space between the vertebral spine processes. This reduces the force required for removal, thus reducing the risk of fracturing the catheter [8]. If catheter breakage occurs, it is essential to perform a CT scan of the spine to determine the track and position of the retained fragment. The decision to remove the retained portion should be made on a case by case basis [16].

In our institution, it is also recommended that a full coagulation profile be performed prior to lumbar drain removal. For patients on a heparin infusion, it should be discontinued 2-4 hours prior to removal and not restarted for a full 12 hours afterward. It is prudent to wait 6-8 hours to start prophylactic once daily dosing and the catheter can be removed 12 hours after the last dose of once daily low molecular weight heparin. Twice daily dosing of low molecular weight heparin (LMWH) is associated with increased risk of spinal hematoma. The first dose of LMWH should be administered no earlier than 24 hours postoperatively. Indwelling catheters should be removed prior to initiation of LMWH thromboprophylaxis. Adequate surgical hemostasis and reversal of coagulopathy should be assured [9].

The optimal management of lumbar drain catheters for TEVAR lacks consensus and requires further evaluation. Given the number of studies revealing benefit and rare incidence of complications, the procedure will continue to be an integral aspect of perioperative care of these patients. Vigilance is required to evaluate the patient not only for neurologic sequelae of their operation, but for potentially catastrophic consequences of the indwelling drain. Through the use of education and awareness at all levels of care, reduction in paraplegia can be safely achieved.

References

