

# Post-dural puncture headache

Ahmed Ghaleb<sup>1</sup>  
Arjang Khorasani<sup>2</sup>  
Devanand Mangar<sup>3</sup>

<sup>1</sup>Department of Anesthesiology,  
University of Arkansas for  
Medical Sciences, Little Rock, AR,

<sup>2</sup>Anesthesiology Residency Program,  
Advocate Illinois Masonic Medical  
Center, Chicago, IL, <sup>3</sup>Florida Gulf to  
Bay Anesthesiology, Tampa General  
Hospital, Tampa, FL, USA

**Abstract:** Since August Bier reported the first case in 1898, post-dural puncture headache (PDPH) has been a problem for patients following dural punctures. Clinical and laboratory research over the last 30 years has shown that use of smaller-gauge needles, particularly of the pencil-point design, are associated with a lower risk of PDPH than traditional cutting point needle tips (Quincke-point needle). A careful history can rule out other causes of headache. A postural component of headache is the sine qua non of PDPH. In high-risk patients < 50 years, post-partum, in the event a large-gauge needle puncture is initiated, an epidural blood patch should be performed within 24–48 hours of dural puncture. The optimum volume of blood has been shown to be 12–20 mL for adult patients. Complications caused by autologous epidural blood patching (AEBP) are rare.

**Keywords:** post-dural puncture headache, gauge, needles, cause, risk, incidence

## Introduction

Post-dural puncture headache (PDPH) is an important iatrogenic cause of patient morbidity in spinal anesthesia, pain management after attempted epidural blocks and after spinal taps. The incidence of dural puncture ranges from 0.16%–1.3% in experienced hands.<sup>1</sup> Post-dural puncture headache develops in 16%–86% of cases after attempted epidural block with large bore needles.<sup>2</sup>

Any breach in the dura may result in PDPH. A breach can be either iatrogenic or spontaneous. Performing an epidural or spinal anesthetic or a diagnostic myelogram can produce the very distinctive PDPH. It can occur immediately or as long as 48 hours post-procedure.

Spontaneous intrathecal CSF leaks (SIH) leading to headache are rare, with a prevalence of approximately 1:50,000 persons, and are more common in women, with a female-male ratio of 3:1. Spontaneous intracranial hypotension is usually seen in the cervical-thoracic region and is also associated with co-morbidities like Marfan's syndrome, neurofibromatosis, connective tissue disorders and Ehlers–Danlos syndrome.

The International Headache Society recognizes the absence of a definitive pain description for the headache associated with SIH. In its criteria of 1998, the International Headache Society classify low CSF pressure, which includes CSF fistula headache: “Posttraumatic, postoperative or idiopathic fluid leak demonstrated by measurement of glucose concentration in leaking fluid, or by leakage of spinally injected dye or radioactive tracer” with characteristics of post-lumbar puncture headache.

Correspondence: Ahmed Ghaleb  
University of Arkansas for Medical  
Sciences, 4301 West Markham,  
Slot 515, Little Rock, AR 72205, USA  
Tel +1 501 686 8818  
Fax +1 501 526 4216  
Email ghalebahmed@uams.edu

According to the International Headache Society, the criteria for a low CSF pressure PDPH<sup>3,4</sup> include a headache that develops less than 7 days after a spinal puncture, occurs or worsens less than 15 minutes after assuming the upright position and improves after less than 30 minutes in the recumbent position. The headache should disappear within 14 days after a spinal puncture; if it persists it is called a CSF fistula headache.

## History

The history of spinal anesthesia can be traced back to the late 1800s when Wynter and Quincke aspirated cerebrospinal fluid from patients with tuberculous meningitis in an attempt to lower intracranial pressure.<sup>5</sup> August Bier performed spinal on himself and eight other subjects using 10–15 mg of cocaine. Four of the nine people, including Professor Bier, developed PDPH.<sup>6</sup>

## Anatomy and pathophysiology

Anatomically, the spinal dura mater extends from the foramen magnum to the second segment of the sacrum. It consists of a dense connective tissue matrix of collagen and elastic fibers. The average adult produces about 500 mL of CSF per day, or 21 mL per hour (0.3 mL/kg/hr), with 90% coming from the choroid plexus, and 10% from the brain substance itself. A total of about 150 mL of CSF circulates at any one time and is absorbed by the arachnoid villi. The cause of PDPH is not entirely certain. The best explanation is that low CSF pressure results from CSF leakage through a dural or arachnoid tear; a leakage that exceeds the rate of CSF production.<sup>4</sup> As little as 10% loss of CSF volume can cause an orthostatic headache. There are two basic theoretical mechanisms to explain PDPH. One is reflex vasodilatation of the meningeal vessels due to the lowered CSF pressure. The other is traction on the pain-sensitive intracranial structures in the upright position. Traction on the upper cervical nerves including C1, C2, and C3, causes pain in the neck and shoulders. Traction on the fifth cranial nerve causes a frontal headache. Traction on the sixth cranial nerve causes visual symptoms. Pain in the occipital region is due to the traction of the ninth and tenth cranial nerves.

## Needle size and incidence of PDPH

The incidence of PDPH is directly related to the needle diameter that pierces the dura mater.<sup>7</sup>

Diagnostic lumbar puncture (LP) requires 22-gauge needles to facilitate measurement of opening pressure and withdrawal of CSF over a reasonably brief time period.

With needles smaller than size 22-gauge, collection of 2 mL of CSF may take 6 minutes or longer and measurement of CSF pressure may be less accurate.<sup>8</sup> An addendum to the American Academy of Neurology (AAN) practice guidelines advocated the use of 22-gauge needles, but reported a case series where 25-gauge needles were used successfully.<sup>9</sup>

Although smaller diameter needle punctures used for subarachnoid block decrease the risk of PDPH, these needles are technically difficult to use and are associated with a lower success rate in spinal anesthesia,<sup>10</sup> especially in inexperienced hands. This is due to failure to recognize dural puncture secondary to slow flow through a small needle, leading to multiple and repeated puncture attempts. The incidence of PDPH with the 25-gauge Whitacre (non-cutting) needle is less than with the 27-gauge Quincke (cutting) needle.<sup>7</sup> Morbidity associated with lumbar puncture can be decreased by the proper selection of an appropriate needle gauge and needle tip configuration.<sup>11,12</sup>

## Direction of bevel

Spinal needles are designed as cutting bevels, as in the Quincke-type, or pencil point, as in the Whitacre-type spinal needle. Dural fibers were once believed to run longitudinally;<sup>13</sup> however microscopic dissection of the dura mater in cadavers revealed that dural fibers do not run longitudinally or in a parallel fashion. The dura is a laminated structure built up from well-defined layers oriented concentrically around the medulla spinalis.<sup>14</sup> Orienting the bevel of a cutting needle probably needs further consideration before making absolute, blanket statements regarding the etiology of dural puncture leaks. The use of a paramedian approach to the subarachnoid space has been suggested as a means of reducing PDPH particularly when using cutting needles.<sup>15</sup>

Electron microscopy has shown that pencil point needles are more traumatic to the dura than the cut bevel needles. It is postulated that a pencil point needle produces an irregular tear in the dura and the subsequent inflammatory reaction reduces CSF leakage more effectively than the clean U-shaped puncture seen with a cutting-bevel needle, which decreases the risk of PDPH.<sup>16</sup>

## Dura mater and response to trauma

After perforation of the dura there will be leakage of CSF. In neurosurgical experience even minor perforations need to be closed, either directly or through the application of synthetic or biological dural graft material. Failure to close the dural perforation may lead to adhesions, continuing CSF

leak, and the risk of infection. It was thought that the closure was facilitated through fibroblastic proliferation from the cut edge of the dura. Work published in 1959<sup>17</sup> dismissed the notion that the fibroblastic proliferation arose from the cut edge of the dura. This study maintained that the dural repair was facilitated by fibroblastic proliferation from surrounding tissue and blood clots. The study also noted that dural repair was promoted by damage to the pia arachnoid, the underlying brain and the presence of blood clots. It is therefore possible that a spinal needle carefully placed in the subarachnoid space does not promote dural healing, as trauma to adjacent tissue is minimal. Indeed, the observation that blood promotes dural healing agrees with Gormley's original observation that bloody taps were less likely to lead to a post-dural puncture headache as a consequence of a persistent CSF leak.<sup>18</sup>

## Symptoms

PDPH typically manifests as a postural, frontal, frontotemporal, or occipital headache, worsened by ambulation and improved by assuming the decubitus position, occurring within 48 hours after dural puncture. The accompanying symptoms are usually nausea, vomiting and neck stiffness.<sup>1,19</sup> Atypical symptoms after accidental dural puncture have been infrequently described.

Other nonspecific symptoms may occur such as nausea, vomiting, ocular complaints such as photophobia and diplopia, and auditory complaints like tinnitus and hyperacusis. The first case of diplopia after dural puncture was reported by Quincke more than 100 years ago.<sup>19</sup> Diplopia or extraocular muscle paralysis (EOM) after dural puncture has been reported occasionally, primarily in the neurology and ophthalmology literature. Because there seems to be a window period before diplopia manifests after dural puncture, the patient and physician may not always believe that the symptom is secondary to dural puncture, particularly when it occurs after resolution of PDPH. Diplopia usually occurs 4–10 days after dural puncture but can manifest as late as 3 weeks. Full recovery can generally be expected in 2 weeks to 8 months, although permanent cases have rarely been reported.<sup>20</sup>

## Factors influencing incidence

Women, particularly during pregnancy and especially after vaginal delivery, are considered at increased risk for PDPH. The incidence of PDPH is highest between 18 and 30 years of age and declines in children younger than 13 years and adults older than 60 years. The incidence is greater in patients with lower body mass index.<sup>2</sup> Women who are obese or

morbidly obese may actually have a decreased incidence of PDPH. The decreased incidence is due to the increase in intra-abdominal pressure which may act as an abdominal binder helping to seal the defect in the dura and decreasing the loss of CSF. Younger women may be at a greater risk because of increased dural fiber elasticity that maintains a patent dural defect compared to a less elastic dura in older patients.<sup>4</sup> Patients with a headache before lumbar puncture and a prior history of PDPH are also at increased risk. There is no known relationship between the diagnosis of migraine headaches and increased incidence of PDPH after regional anesthesia.<sup>21</sup> There may be some correlation between a history of motion sickness and PDPH. Another important factor is the experience of the person doing the procedure leading to the puncture of the dura. Continuous spinal infusion reduced the incidence of PDPH when compared to single shot spinal, at least according to the study.<sup>22</sup>

## Differential diagnosis

A comprehensive history and physical exam must be carried out before making the diagnosis of PDPH. Spinal abscess, spinal hematoma, septic or aseptic meningitis, intracranial mass lesion, cerebral aneurysm, cerebral edema, myofascial syndrome, arachnoiditis caused by intrathecal steroids, transient neurologic syndrome or related symptoms, unspecific post-dural puncture lumbalgia, neural toxicity of the drugs, and anterior spinal artery syndrome, post-partum cerebral angiopathy and cerebral thrombophlebitis should all be ruled out.<sup>10,23,24</sup> Additional tests such as a cerebral CT scan, or magnetic resonance imaging could be performed in cases with atypical post-dural puncture symptoms, to exclude the possibility of developing serious complications.<sup>25</sup>

Few cases of atypical post-dural puncture symptoms have been reported in the literature. Lybecker et al cited interscapular pain as a "related musculoskeletal symptom," however; no instances of upper back pain are cited among the 75 cases of PDPH reported by the authors.<sup>19</sup> McGrady and Freshwater reported a case of posterior neck pain without headache after spinal anesthesia.<sup>26</sup> Schabel et al reported a case of arm pain with dysesthesia after an unintended dural puncture, and explained it as irritation of the C5 and C6 nerve roots caused by central traction.<sup>24</sup>

## Treatment

### Conservative/symptomatic therapy

The treating clinician must provide emotional support and reassurance to patients with PDPH. Bed rest has been advocated in cases of dural puncture by some clinicians.

However, a recent meta-analysis failed to show that bed rest after dural puncture was better than immediate mobilization in reducing the incidence of PDPH.<sup>27</sup> Bed rest can be associated with a higher incidence of PDPH in particular patient groups.<sup>28</sup> Bed rest may postpone the occurrence of the headache but does not prevent it.

## Pharmacotherapy

### Oral and intravenous medications

Oral hydration remains a popular therapy for PDPH, but there is no evidence that vigorous hydration has any therapeutic benefit, or that it encourages an increased production of cerebrospinal fluid. However, no patient with PDPH should be allowed to become dehydrated.

The efficacy of oral caffeine for the treatment of PDPH was evaluated in 40 postpartum patients.<sup>29</sup> A single oral dose was demonstrated to be safe, effective and should be considered in the early treatment of mild PDPH. Caffeine sodium benzoate, as an intravenous bolus or an infusion can be used to treat PDPH. Caffeine was 75% to 80% effective in the initial treatment of PDPH; however, follow-up 48 hours later revealed that all patients had a return of their headache.<sup>30</sup>

Cosyntropin, a synthetic form of adrenocorticotropic hormone, has been used in the treatment of refractory PDPH. Adrenocorticotropic hormone is believed to work by stimulating the adrenal gland to increase CSF production and  $\beta$ -endorphin output. Caution should be used in patients with diabetes.<sup>31</sup>

The serotonin type 1-d receptor agonist sumatriptan is effective in the treatment of PDPH, with complete resolution of symptoms.<sup>32,33</sup> The drug is expensive, and side effects include pain at the site of injection and chest tightness. Caution must be used in treating patients with ischemic heart disease using sumatriptan.<sup>32,33</sup> Controlled trials are needed to further evaluate the use of sumatriptan for PDPH.

A trend away from conservative management to the use of the blood patch has appeared in recent years. This is based on the relative ineffectiveness of the conservative treatment. For example, over 80% of postpartum patients who are conservatively treated will still have a headache after 1 week.

### Epidural injections

The autologous epidural blood patch (AEBP) has become the 'gold standard' in the treatment of PDPH. As there is some risk of infection when injecting blood into the epidural space, we will discuss the efficacy of some other aqueous agents that have been injected into the epidural space to treat PDPH. Prior to considering the use of epidural injections

of blood or other substances to relieve the symptoms of PDPH, there needs to be a clearly negative history of sepsis and coagulopathy. HIV infection is not considered to be a contraindication to AEBP.

Dextran and 0.9% NaCl (saline) injections into the epidural space transiently increase pressure in the epidural space, which subsequently decreases the leakage of CSF and restores subarachnoid pressure.<sup>34–36</sup> Not only is the success rate moderate, but also anaphylaxis has been reported following the use of dextran for this purpose.<sup>35</sup> Epidural patching with non-blood substances, eg, saline or colloid, are ineffective for prolonged relief,<sup>37</sup> although other substances such as fibrin glue have been used.<sup>38</sup> The utility of epidural morphine was investigated in one RCT.<sup>39,40,41</sup> Epidural morphine 3 mg was given after the end of anaesthesia and another 3 mg was given on the following day. This reduced the incidence of PDPH from 48% (17/25) to 12% (2/25), which translates to a statistically significant reduction in the relative risk of 0.25 (0.08–0.73). There was no respiratory depression, but nausea was numerically more frequent in the morphine group (44% versus 10%,  $P = 0.06$ ).

### Autologous epidural blood patch (AEBP)

The AEBP was first described by Gormley in 1960 for use in PDPH and was later popularized by DiGiovanni et al.<sup>38,42</sup> The proposed mechanism of action of AEBP is tamponade of the dural leakage while simultaneously raising the subarachnoid pressure. Elevation of subarachnoid and epidural pressures remains for only about 20 minutes.<sup>42</sup> MRI evidence confirms a mass effect after injection of epidural blood, with gradual resolution over about 7 hours. Unlike saline, dextran or other fluids, blood is not removed quickly from the epidural space,<sup>43</sup> and it potentially exerts a tamponade effect for much longer periods of time. The autologous blood is thought to form a fibrin clot over the dural rent, allowing CSF volume and hence pressure to normalize as new CSF is generated.<sup>44</sup>

Abouleish et al, summarized 524 cases of AEBP reported by eleven centers.<sup>45</sup> Persistent symptomatic relief of PDPH following epidural blood patch was >95%, particularly when using volumes of blood >15 mL. In this review, using volumes of blood greater than 20 mL offered no advantages, as it is known that 20 mL spreads about 9–10 spinal segments when administered to patients in the sitting position.<sup>12</sup>

Some studies have demonstrated lower success rates, with only 61%–75% of patients demonstrating sustained benefit. These lower success rates may reflect dural puncture occurring with large-bore epidural needles versus smaller-gauge spinal needles.<sup>44,46,47</sup> In obstetrical studies, the success rate

of epidural blood patching for PDPH is lower because the dural hole made by 18 gauge Tuohy needles results in a large leakage of CSF, necessitating a second blood patch in as many as 29% of patients.<sup>34,47</sup>

### The technique of AEBP

The procedure is performed only after a careful history to exclude other causes of headache. While some authors have recommended the administration of prophylactic antibiotics for the procedure, they are generally not used. Rarely, if fluoroscopy is being utilized, the prone position may be selected. The preferred interspace for injection is the one below the previous injection site, because blood preferentially rises cephalad following its injection into the lumbar epidural space.<sup>43,48</sup> Usually, at least 20 mL of blood are aseptically withdrawn. In children, 0.2–0.3 mL/kg of blood is needed. Phlebotomy should be attempted after first identifying the epidural space to avoid clotting. The blood is carefully, and aseptically, transferred to the anesthesiologist, who injects it slowly through the epidural needle until one of the following endpoints occurs: (a) complaint of back pain, neck pain or radicular pain in the leg or worsening headache during the performance of the epidural injection, or (b) once at least 20 mL have been successfully injected without complaint by the patient. The patient is advised to avoid straining, bending or heavy lifting for 2–3 days to allow the dural hole to heal.

In regard to the optimum volume of autologous blood to be injected epidurally, Taivainen et al found that using 20 mL standard in all patients was equivalent to 1–15 mL variably administered based upon height.<sup>46</sup> Others have advocated more generous volumes. Crawford found that 20 mL was associated with 96% success versus 70% success using 6–15 mL.<sup>51</sup> The ideal time to perform an epidural blood patch is within 24 hours of puncture.<sup>50</sup> Treatment failure after blood patching may reflect continued transdural leak;<sup>51</sup> in this case, the blood patch should be repeated while keeping the patient flat for 24 hours afterwards to reduce the flow of CSF through the dural rent.

Complications following AEBP include the following: backache (35%), neck pain (0.9%), and transient temperature elevations (5%) lasting 24–48 hours. Bleeding, infection, repeat dural puncture, and arachnoiditis from blood injected into the subarachnoid space have been reported. There have been at least two cases of facial nerve paralysis reported following autologous blood patch, both of which resolved spontaneously. Lowe and McCullough suggested that the etiology is ischemia of the 7th nerve resulting from decreased blood supply after an increase in intracranial pressure due

to the injection of blood in the epidural space.<sup>52</sup> There has also been at least one case of intractable dizziness, vertigo, tinnitus and ataxia.<sup>53,54</sup> Blood patching has occasionally been associated with vasovagal syncope.<sup>55</sup>

### Prophylactic blood patch

Some have suggested that blood patching be performed as a prophylactic measure (ie, prior to the development of a headache) in cases of unintended dural puncture occurring after the insertion of 17–18 gauge epidural needle into the subarachnoid space, particularly when there has been loss of considerable quantities of CSF. To date, there have been no large, prospective studies to advocate this practice, although some anesthesia practitioners still use it. Existing studies regarding prophylactic epidural blood patch are limited by small patient numbers.<sup>56</sup> Also, there have been limitations suggested by the small volumes of blood injected in some reviews.<sup>58</sup> If one chooses to perform prophylactic AEBP, some caveats are in order; one should avoid prophylactic AEBP immediately following local anesthetic (LA) epidural top-off dose administration, because the resultant high epidural pressure has resulted in at least one case of total spinal block.<sup>59</sup> Also, the presence of LA in the epidural space may theoretically interfere with subsequent blood clot formation.<sup>60</sup> Some are strongly advocated for prophylactic blood patch, particularly if an epidural catheter is in place, they argue that it avoids the need for another epidural puncture, even though strong clinical evidence is lacking.<sup>37,61,62</sup>

### Prognosis

Since PDPH is usually a self-limited condition, the prognosis is not significantly affected by treatment. The majority of headaches resolve within a week with conservative management (rest, hydration, symptomatic treatment).<sup>63</sup> In an older study, 53% of headaches resolve in 4 days, 72% in 7 days, and 85% within 6 weeks.<sup>63</sup> In a small minority of cases symptoms may persist for weeks, months, or even years.<sup>64</sup> Rarely, the headache can become chronic with the longest reported headache after LP lasting 5 years.<sup>47,48,50,51</sup> As PDPH begins to improve, headache severity decreases, patient mobility increases, and it takes longer for headache following postural changes to develop.<sup>52</sup> It is unclear if patients with chronic headache or pain disorders experience more severe and/or prolonged PDPH.

### Conclusion

Although not life-threatening, PDPH carries substantial morbidity by restricting activities of daily life. Current

noninvasive treatments, including bed rest, fluids, analgesics, caffeine, and sumatriptan, only temporize the discomfort.<sup>33</sup> Epidural blood patch remains the invasive treatment of choice, with approximately 70% prolonged success after initial injection.<sup>65</sup> The benefit of prophylactic blood patching is not so clear but deserves consideration in those most at risk from a headache, such as the parturient, and after accidental dural perforation with a Tuohy needle. Surgical closure of the dural tear remains an option of last resort.

## Disclosure

The author reports no conflict of interest in this work.

## References

- Reynolds F. Dural puncture and headache. *BMJ*. 1993;306(6882): 874–876.
- Liu S, Carpenter RL, Neal JM. Epidural anesthesia and analgesia. Their role in postoperative outcome. *Anesthesiology*. 1995;82(6): 1474–1506.
- Diaz JH. Epidemiology and outcome of postural headache management in spontaneous intracranial hypotension. *Reg Anesth Pain Med*. 2001;26(6):582–587.
- Evans RW. Complications of lumbar puncture. *Neurol Clin*. 1998;16(1): 83–105.
- Harrington BE. Post-dural puncture headache and the development of the epidural blood patch. *Reg Anesth Pain Med*. 2004;29(2): 136–163.
- Turnbull DK, Shepherd DB. Post-dural puncture headache: Pathogenesis, prevention and treatment. *Br J Anaesth*. 2003;91(5):718–729.
- Lambert DH, Hurley RJ, Hertwig L, Datta S. Role of needle gauge and tip configuration in the production of lumbar puncture headache. *Br J Anaesth*. 1997;22(1):66–72.
- Strachan A, Train J. Lumbar puncture and headache. Aspirating cerebrospinal fluid speeds up procedure. *Br Med J*. 1998;317(7136): 1018–1019.
- Armon C, Evans RW. Addendum to assessment: Prevention of post-lumbar puncture headaches. Report of the Therapeutics and Technology Assessment Subcommittee of the American Academy of Neurology. *Neurology*. 2005;65(4):510–512.
- Horlocker TT. Complications of spinal and epidural anesthesia. *Anesthesiol Clin North America*. 2000;50(2):461–485.
- Vandam LD, Dripps RD. Long-term follow-up of patients who received 10,098 spinal anesthetics; syndrome of decreased intracranial pressure (headache and ocular and auditory difficulties). *J Am Med Assoc*. 1956;161(7):586–590.
- Gielen M. Post dural puncture headache: a review. *Reg Anesth*. 1989; 14(3):101–106.
- Norris MC, Leighton BL, Desimone CA. Needle bevel direction and headache after inadvertent dural puncture. *Anesthesiology*. 1989; 70(5):729–731.
- Reina MA, Dittmann M, Lopez Garcia A, van Zundert A. New perspectives in the microscopic structure of human dura mater in the dorsolumbar region. *Reg Anesth*. 1997;22(2):161–166.
- Ready LB, Cuplin S, Haschke RH, Nessly M. Spinal needle determinants of rate of transdural fluid leakage. *Anaesth Analg*. 1989; 69(4):457–460.
- Reina MA, de Leon-Casasola OA, Lopez A, De Andres J, Martin S, Mora M. An in vitro study of dural lesions produced by 25-gauge Quincke and Whitacre needles evaluated by scanning electron microscopy. *Reg Anesth Pain Med*. 2000;25(4):393–402.
- Keener EB. An experimental study of reactions of the dura mater to wounding and loss of substance. *J Neurosurg*. 1959;16(4):424–447.
- Gormley JB. Treatment of post-spinal headache. *Anesthesiology*. 1960; 21:565–566.
- Lybecker H, Djernes M, Schmidt JF. Post-dural puncture headache PDPH: onset, duration, severity, and associated symptoms. An analysis of 75 consecutive patients with PDPH. *Acta Anaesthesiol Scand*. 1995; 39(5):605–612.
- Nishio I, Williams BA, Williams JP. Diplopia: a complication of dural puncture. *Anesthesiology*. 2004;100(1):158–164.
- Bader AM. Neurologic and neuromuscular disease in the obstetric patient. *Anesthesiol Clin North America*. 1998;16:459–476.
- Maurer K, Bonvini JM, EkatoDRAMIS G, Serena S, Borgeat A. Continuous spinal anesthesia/analgesia vs single-shot spinal anesthesia with patient-controlled analgesia for elective hip arthroplasty. *Acta Anaesthesiol Scand*. 2003;47(7):878–883.
- Abram SE. Treatment of lumbosacral radiculopathy with epidural steroids. *Anesthesiology*. 1999;91(6):1937–1941.
- Errando CL. Transient neurologic syndrome, transient radicular irritation, or postspinal musculoskeletal symptoms: Are we describing the same “syndrome” in all patients? *Reg Anesth Pain Med*. 2001; 26(2):178–180.
- Mokri B, Atkinson JLD, Dorek DW, et al. Syndrome of orthostatic headaches and diffuse pachymeningeal gadolinium enhancement. *Mayo Clin Proc*. 1997;72(5):400–404.
- McGrady EM, Freshwater JV. Spinal headache – with no headache. *Anaesthesia*. 1997;46(9):707.
- Thoenissen J, Beckers H, Lamy W, Domanovits H, Laggner A, Müllner M. Does bed rest after cervical or lumbar puncture prevent headache? A systematic review and meta-analysis. *CMAJ*. 2001; 165(10):1311–1316.
- Fanoulaki A, Siantopoulos C, Andreopoulos K. Is early mobilization associated with lower incidence of postspinal headache? A controlled trial in 69 urologic patients. *Anesthesiol Reanim*. 1991; 16(6):375–378.
- Thornberry EA, Thomas TA. Posture and post-spinal headache. A controlled trial in 80 obstetric patients. *Br J Anaesth*. 1988;60(2): 195–197.
- Camann WR, Murray RS, Mushlin PS, Lambert DH. Effects of oral caffeine on post-dural puncture headache: a double-blind, placebo-controlled trial. *Anesth Analg*. 1990;70(2):181–184.
- Carter BL, Pasupuleti R. Use of intravenous cosyntropin in the treatment of post-dural puncture headache. *Anesthesiology*. 2000;92(1): 272–274.
- Collier BB. Treatment for post dural puncture headache. *Br J Anaesth*. 1994;72(3):366–367.
- Carp H, Singh PJ, Vadhera R, Jayaram A. Effects of the serotonin-receptor agonist sumatriptan on post-dural puncture headache: report of six cases. *Anesth Analg*. 1994;79(1):180–182.
- Hodgson C, Roitberg-Henry A. The use of sumatriptan in the treatment of post-dural puncture headache. *Anaesthesia*. 1997;52(8):808.
- Stride PC, Cooper PC. Dural taps revisited. A 20-year survey from Birmingham Maternity Hospital. *Anaesthesia*. 1993;48(3): 247–255.
- Trivedi NS, Eddi D, Shevde K. Headache prevention following accidental dural puncture in obstetric patients. *J Clin Anesth*. 1993;5(1): 42–45.
- Barrios-Alcaron J, Aldrete JA, Paragas-Tapia D. Relief of post-lumbar puncture headache with epidural dextran 40: a preliminary report. *Reg Anesth*. 1989;14(2):78–80.
- DiGiovanni AJ, Dunbar BS. Epidural injections of autologous blood for postlumbar puncture headache. *Anesth Analg*. 1970;49(2):268–271.
- Duffy PJ, Crosby ET. The epidural blood patch: Resolving the controversies. *Can J Anaesth*. 1999;46(9):878–886.
- Al-Metwalli RR. Epidural morphine injections for prevention of post dural puncture headache. *Anaesthesia*. 2008;63(8):847–850.
- Crul BJ, Gerritse BM, van Dongen RT, Schoonderwaldt HC. Epidural fibrin glue injection stops persistent post-dural puncture headache. *Anesthesiology*. 1999;91(2):576–577.

42. DiGiovanni AJ, Galbert MW, Wahle WM. Epidural injection of autologous blood for postlumbar-puncture headache. II: Additional clinical experiences and laboratory investigation. *Anesth Analg.* 1972;51(2):226–232.
43. Beards SC, Jackson A, Griffiths AG, Horsman EL. Magnetic resonance imaging of extradural blood patches: appearances from 30 min to 18 h. *Br J Anaesth.* 1993;71(2):182–188.
44. Szeinfeld M, Ihmeidan IH, Moser MM, Machado R, Klose KJ, Serafini AN. Epidural blood patch: evaluation of the volume and spread of blood injected into the epidural space. *Anesthesiology.* 1986;64(6):820–822.
45. Abouleish E, Vega S, Blendinger I, Tio TO. Long-term follow-up of epidural blood patch. *Anesth Analg.* 1975;54(4):459–463.
46. Taivainen T, Pitkanen M, Tuominen M, Rosenberg PH. Efficacy of epidural blood patch for post-dural puncture headache. *Acta Anaesthesiol Scand.* 1993;37(7):702–705.
47. Safa-Tisseront V, Thormann F, Malassine P, et al. Effectiveness of epidural blood patch in the management of post-dural puncture headache. *Anesthesiology.* 2001;95(2):334–339.
48. Djurjuus HJ, Rasmussen M, Jensen EH. Epidural blood patch illustrated by CT-epidurography. *Acta Anaesthesiol Scand.* 1995;39(5):613–617.
49. Rosenberg PH, Heavner JE. In vitro study of the effect of epidural blood patch on leakage through a dural puncture. *Anesth Analg.* 1985;64(5):501–504.
50. Loeser EA, Hill GE, Bennett GM, Sederberg JH. Time vs success rate for epidural blood patch. *Anesthesiology.* 1978;49(2):147–148.
51. Crawford JS. Experiences with epidural blood patch. *Anaesthesia.* 1980;35(5):513–515.
52. Lowe DM, McCullough AM. 7th nerve palsy after extradural blood patch. *Br J Anaesth.* 1990;65(5):721–722.
53. Ostheimer GW, Palahiuk RJ, Shnider SM. Epidural blood patch for post-lumbar puncture headache. *Anesthesiology.* 1974;41(3):307–308.
54. Walpole JB. Blood patch for spinal headache. A recurrence and a complication. *Anaesthesia.* 1975;30(6):783–785.
55. Andrews PJ, Ackerman WE, Juneja M, Cases-Cristobal V, Rigor BML. Transient bradycardia associated with extradural blood patch after inadvertent dural puncture in parturients. *Br J Anaesth.* 1992;69(4):401–403.
56. Colonna-Romano P, Shapiro BE. Unintentional dural puncture and prophylactic epidural blood patch in obstetrics. *Anaesth Analg.* 1989;69(4):522–523.
57. Quaynor H, Corbey M. Extradural blood patch – why delay? *Br J Anaesth.* 1985;57(5):538–540.
58. Palahniuk RJ, Cumming M. Prophylactic blood patch does not prevent post lumbar puncture headache. *Can Anaesth Soc J.* 1979;26(2):132–133.
59. Leivers D. Total spinal anesthesia following early prophylactic epidural blood patch. *Anesthesiology.* 1990;73(6):1287–1289.
60. Tobias MD, Pilla MA, Rogers C, Jobes DR. Lidocaine inhibits blood coagulation: implications for epidural blood patch. *Anesth Analg.* 1996;82(4):766–769.
61. Stevens RA. Neuraxis blocks. In Brown DL, editor. *Regional Anaesthesia and Analgesia.* Philadelphia: WB Saunders; 1996:352.
62. Cheek TG, Banner R, Sauster J, Gache BB. Prophylactic extradural blood patch is effective. A preliminary communication. *Br J Anaesth.* 1988;61(3):340–342.
63. Dripps RD, Vandam LD. Long term follow up of patients who received 10,098 spinal anesthetics: failed to discover major neurological sequelae. *J Am Med Assoc.* 1954;156(6):1486–1491.
64. Lybecker H, Djerfås J, Schmitt J. Post-dural puncture headache (PDPH): onset, duration, severity, and associated symptoms. An analysis of 76 consecutive patients with PDPH. *Acta Anaesthesiol Scand.* 1995;39(5):607–612.
65. Vellaria SB, Thomas PS, Rosenbaum AE, Wasenko JJ, Fellows DG. Magnetic resonance imaging of cerebrospinal fluid leak and tamponade effect of blood patch in post-dural puncture headache. *Anesth Analg.* 1997;84(3):585–590.

RETRACTED

### International Journal of General Medicine

### Publish your work in this journal

The International Journal of General Medicine is an international, peer-reviewed open-access journal that focuses on general and internal medicine, pathogenesis, epidemiology, diagnosis, monitoring and treatment protocols. The journal is characterized by the rapid reporting of reviews, original research and clinical studies across all disease areas.

Submit your manuscript here: <http://www.dovepress.com/international-journal-of-general-medicine-journal>

Dovepress

A key focus is the elucidation of disease processes and management protocols resulting in improved outcomes for the patient. The manuscript management system is completely online and includes a very quick and fair peer-review system. Visit <http://www.dovepress.com/testimonials.php> to read real quotes from published authors.