

Complications in spinal anaesthesia

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Introduc on

Spinal anaesthesia is one of the most popular and widely used anaesthe c procedures. It is a simple, cost e ec ve and e cient technique that provides complete sensory and motor block, as well as postopera ve analgaesia with a high success rate. Several advantages of spinal anaesthesia include a decreased incidence of deep vein thrombosis, reduced intraopera ve blood loss, as well as the preven on of pulmonary aspira on in case of emergency, especiallyin pa ents with poten al airway problems and known respiratory diseases.

Due to the invasive nature of spinal anaesthesia, there are several types of complica ons that may occur with di erent incidence. At least some of these problems appear to be inevitable and as such, it is not possible to eliminate them all. Fortunately, more severe neurological complica ons such as death, neuropathy, arachnoidi s and permanent neurologic injury are seldom observed. In a na onal survey performed in the UK, the incidence of permanent neurologic injury and death ranged from 0. 7 to 1. 8 in 100, 000 pa ents¹. On the other hand, proper pa ent selec on, me culous a en on to detail, well-known pa ent related changes and in the case of di cult circumstances, using image techniques [x rays, fluoroscopy and ultrasound] as a guide may help to prevent or decrease complica ons.

Increasing co-morbidi es, concomitant medica on, surgery for advanced malignancy, pa ents with compromised immune systems, as well as instances of infec on poses a real challenge to the use of spinal anaesthesia. Pa ents with degenera ve vertebral anomalies or who have undergone previous spinal surgeries are also di cult cases; these require further evalua on and an increase in e orts for properly performing intrathecal anaesthesia and analgaesia in contextswhere it may lead to undesirable consequences.

anaesthesia that occurs when the sympathe c chain becomes blocked, especially when higher dermatome levels are needed. A drop in blood pressure may ini ate nausea and vomi ng, indica ng ischaemia on the spinal cord, which in turn induces an undesired condi on for the pa ent and opera ng sta. Blood pressure changes between the le lateral to supine posi on has been determined as an indicator for predic ng a periopera ve decrease in obstetric pa ents undergoing caesarean delivery under spinal anaesthesia².

In a non-obstetric study popula on, changing pa ents to the Trendelenburg posi on for 10 minutes immediately following a spinal block has been demonstrated as e cient, as has loading with a lactated ringer or 6% hydroxyl ethyl starch solu on by means of maintaining cardiac output. Co-hydra on is more e cient than prehydra on and colloid loading is be er for maintaining cardiac output and blood pressure ³. In their report, Shin et al. ⁴ inves gated the influence of crystalloid and colloid loading on cerebrospinal fluid movements in volunteers, as well as the spread of local anaesthe cs in pa ents. Although crystalloid pretreatment delayed the cranial spread of the block, it induced cerebrospinal fluid produc on, which may be valuable in the case of postdural-puncture headache [PDPH].

In case of pregnancy, a decrease in blood pressure at the cri cal level may a ect both motherand baby, and result in more serious outcomes over a longer period. Increased venodilata on under the influence of progesterone or prostaglandins may also contribute to changes in blood pressure. The presence of hypertension, advanced age, increased body massindex, higher birth weight and higher block are considered as risk factors in hypotension performed with spinal anaesthesia. Fluid loading, lateral It or wedge performed under the right bu ock to prevent aortocaval compression, or vasopressor therapy, cons tutes preven ve measurements to treat hypotension in

Hypotension

Hypotension is an inevitable complica on of spinal



obstetric pa ents. The influence of aortocaval pressure or other determinants remain controversial in terms of how they contribute to haemodynamics during spinal anaesthesia. In order to a enuate the e ect of hypotension, the influence of posi oning the pa ent on a lateral decubitus posi on for a brief period was inves gated. The hypotension episode showed a slight delay, but the incidence of hypotension or drug use was the same as observed in pa ents lying supine⁵. In an editorial, Sharwood-Smith and Drummond ⁶ cri cized the role of vena caval compression in light of the presence of persistent vasoconstric on, such as observed in pa ents with pre-edampsia, which is known to be volumen depleted or hypovolemic; hypotension, however, was observed to a lesser extent. These observa ons jus fy vasopressor therapy and indicate that sympathe c block a ec ng arterial vasculature might be a major concern related to arterial pressure drop⁶.

Time, dura on and the selec on of vasoac ve drugs are controversial issues where obstetric pa ents are concerned. Ephedrine may s mulate beta adrenergic receptors by crossing the umbilical cord and increasing foetal acidosis; therefore, phenylephrine has become the vasopressor of choice ⁷. Variable infusion has been demonstrated as being superior to the fixed infusion and 25-50 µg min-1 was demonstrated to be su cient or maintaining hemodynamic status Non-invasive cardiac output monitoring might be indicated for a pa ent with severe cardiac disease ⁸. On the other hand, a phenylephrine infusion may induce bradycardia, presumed (or) indicated by the baroreflex receptor mediated mechanism⁹.

Hypothermia

A decrease in body temperature is commonly encountered a er neuraxial anaesthesia. Subarachnoid local anaesthe c administra on blocks all a erents of skin temperature that pa ents are unable to release the decrease in core temperature. Vasodila on due to sympathe c blockade increases skin blood flow, which allows for lowering the body's core temperatura in a reliable manner. In preparing the skin for surgery with an sep c solu ons, especially when performed on a large area, evapora on from surgical field and irriga on solu ons, or fluid infusion at a higher rate, may also contribute to hypothermia during surgery. [Fig 1] A decrease in core temperature may ini ate shivering, especially during the postopera ve period, which increasesoxygen consump on. Hypothermiaisknown to induce hyper coagula on and infec ons. Special care should be exerted to decrease this physiologic stress, especially in paediatric, obstetric and pa ents in advanced age, since it may lead to serious consequences, including low perfusion to the vital organs, coronary ischaemia and infec on¹⁰.

It is crucial to warm the pa ent with blankets, surgical thermal ma resses, forced air heathers, by using prewarmed irriga on, intravenous solu ons and blood products in order to decrease the severity of this complica on.



Figure 1. Large areas of skin prepped with a Povidone-iodine solu on and uncovered during surgery favour hypothermiaand its complica ons [Source: www.anestesia-dolor.org].

Post-dural-puncture headache

PDPH is a troublesome complica on, mostly observed in middle-aged women and the obstetric popula on. Lower body mass index, previous PDPH and the presence of chronic headaches are other risk factors. Headache rarely occurs in the paediatric popula on, especially in neonates, but some physicians believe that this may be due to the inability to communicate pain in early childhood. PDPH also decreases with age, which may be related to changes in the composi on of cerebral content, which increase on cerebrospinal fluid [CSF] that may compensate and prevents its occurrence. PDPH requires di eren a on from other causes of headache¹¹. It typically occurs in the fronto-occipital region with nuchal rigidity and ini ates when moving from the supine posi on to si ng or standing up. It may vary from mild to severe and the type of pain may be dull, throbbing or burning. Ver go, nausea and vomi ng might be observed due to PDPH in some pa ents. Headache typically appears on the second day following the dural puncture and can range from las ing one to four days, but



may be observed as early as 20 min a er the dural puncture ¹². The leaking of CSF across the dural hole may ini ate PDPH. This is explained by the following mechanisms: a decrease in intracranial pressure causes the trac on of pain sensi ve cranial structures, the deple on of CSF volume may induce compensatory cerebral vasodilata on [the Monroe-Kelly doctrine] and the ac va on of adenosine receptors may cause cerebral vasodilata on ¹³.

The incidence of PDPH has been reported at a level of 2.5% when using a 25 G pencil point needle in obstetric pa ents¹⁴; in the non-obstetric popula on, the incidence of PDPH is as low as 0.37% when using fine spinal needles [15]. Cu ng edge needles are not recommended for spinal anaesthesia, due to the increased incidence of PDPH, even when using fine needles in pa ents undergoing an orectal surgery¹⁶.

Accidental dural entry is a more distressing event that occurs while advancing the Tuohy needle or epidural catheter, resul ng in PDPH at a level of about 75%. The epidural catheter is presumed to introduce from weak points of dura which may occurs with Tuohy needle Incidence of this occurring has been reported as 0.5% in an obstetrical referral centre ¹⁷. It is not possible to recognize or observe clear CSF in needles or catheters in all pa ents. Therefore, as a treatment tool, re-inser ng the epidural catheter in the perforated dura mater with the intent to decrease PDPH does not succeed in all pa ents.

Pa ent posi on when performing spinal anaesthesia, the experience of the physician and using finer needles do not appear to influence the occurrence of PDPH ¹⁷. Pneumocephalus with subsequent PDPH is a rare but well-described complica on of uninten onal dural puncture. It has late dinical onset manifesta ons and can induce a long-las ng headache as a result of accidental dural entry when epidural anaesthesia is performed by means of the loss of resistance technique, using air¹⁸.

The treatment algorithm depends on the severity of PDPH. Conserva ve treatment consists of bed rest and oral or intravenous fluid replacement. Pharmacological therapy includes analgaesics, vasoconstrictors or drugs that increase CSF produc on. Paracetamol or nonsteroidal an -inflammatory drugs are used as first step treatment. Vasoconstrictors, such as ca eine and

Sumatripan, have been used but with limited benefits. Ca eine should be prescribed with cau on due to the pa ent having a lowered convulsion threshold and long term administra on is not advised. Gabapen n has also been used successfully for the treatment of PDPH. Drug therapy may provide relief, but do not completely resolve the symptoms. Epidural morphine has also been demonstrated as beneficial but may leak from the dural hole into the intrathecal space and has well-known side e ectssuch aspruritus, nausea and vomi ng¹⁹.

Although controversies surrounding it remain, the epidural blood patch [EBP] remains the gold standard for trea ng PDPH. A sterile sample of 15-20 mL autologous blood is drawn from the pa ent and immediately injected at the same or a lower level inside the epidural space, un I backache or dullness can be felt. It is generally performed a er wai ng 24 hours following the epidural block. If PDPH persists, a second EBP a week later may be necessary. A third EBP is seldom needed. In a series of cases, the volume of blood needed was reduced while performing EBP under fluoroscopy guidance²⁰.

The witnessed accidental dural entry has di erent treatment op ons Advancing an epidural catheter to the subarachnoid space, injec ng10mLof saline ini ally and leaving the catheter in place for 24 hours are helpful for decreasing the incidence of PDPH. A catheter is believed to induce inflammatory reac on to the dural hole and closure may occur during withdrawal of catheter. At the very least, the presence of the epidural catheter may impede CSF leakage.

Epidural catheter placement in a di erent interspace has also shown poten al benefits. Epidural saline or a dextran infusion for crea ng a fluid column has limited therapeu c e cacy, possibly due to the easy reabsorp on from dural veins ²¹. Fibrin glue was also used, especially in pa ents who refuted the therapy or any other contraindica ons such as coexis ng systemic infec on ²². Surgical treatment is the final step; this only occursif chronic leakage persists²³.

Cranial hypotension and long-las ng CSF loss may distract cerebral bridging veins that can easily rupture and lead to acute or chronic subdural or subarachnoid haematoma occurring. Cau on should be applied when spinal anaesthesia is implemented in a pa ent who has experienced recent cranial trauma, the likes of which may either facilitate or confuse the outcome.



Neurological inves ga on should be performed when the headache lasts more than a few days and is resistant to the conserva ve treatment². Spinal haematoma is a rare event that may relate to the direct needle trauma²⁵.

Chronic leakage and cranial hypotension may influence cranial nerves and nerve palsies may rarely occur due to compression or altered blood supply. It mostly a ects the VI cranial nerve; the reason for this might be a ributed to its longer course (or path) in cranium²⁶. Treatment modali es against CSF leakage and specific therapy for nerve palsy, including cor costeroids, have been demonstrated as being beneficial ²⁷. Altered mental status and speech or stupor may be observed with intracranial hypotension, headache, nausea and vomi ng, and determined as posterior reversible encephalopathy syndrome presen ng as oedema in the posterior cerebral por ons with MRI. This syndrome is largely related to the systemic illness that was first described in an obstetric pa ent following spinal anaesthesia. The late onset and courseof the syndrome implicates compression of posterior por on of the brain vault due to chronicloss of CSF ²⁸. Reversible cerebral vasoconstrictor syndrome is another en ty that has similarclinical features such as headache but lacks imaging findings²⁹.

Hearing lossisanother complica on related to loss of CSF during spinal anaesthesia. Hearing disability especially a ects low frequencies on audiometry and commonly occurs at the second post spinal block. Studies have shown that aims to decrease incidences of leaking CSF usingfine and pencil-point- pped spinal needles can decrease this complica on ³⁰. Type intravenous fluid loading either using crystalloid or colloid do not seems to largely influence or prevents the presence of auditory malfunc ons³¹.

Transient neurologic symptoms

Radicular symptoms, including pain, a burning sensa on on the bu ocks, dysaesthesia and paraesthesia may be observed following spinal anaesthesia. These symptoms generally subside within two days But these clinical features are alarming for possible serious consequences. There is no representa on of these symptoms on radiographs, CT or MRI. Ambulatory surgery, lithotomy posi on, the type of local anaesthe cused, as well as the concentra on of dextrose and osmolarity has being men oned as contribu ng factors for transient neurologic symptoms. The use of spinal lidocaine is one factor that may increase the incidence of transient neurologic symptoms, especially when some factors are combined. An increase in local anaesthe concentra on by pooling and maldistribu on may also increase the incidence of this complica on ³². In a review by Zaricet al. ³³, the authors indicated that the rela ve risk was about seven-to eight-fold lower with other local anaesthe cs such as bupivacaine, mepivacaine, and prilocaine.

Urinary reten on

Bladder distension during the postopera ve period produces discomfort to pa ents and unless relieved, leads to more severe complica ons, including permanent injury to the detrusor muscle. Spinal anaesthesia influences urina on by blocking all a erent nerve fibres, rendering the pa ent unable to feel bladder distension or urinary urgency. Bladder catheteriza on is not innocuous; it carries the risk of urethral trauma and more severe complica ons, including infec on and haematologic spread that may reach the surgical site ³⁴. Urodynamic studies indicate that the func on of the detrusor muscle returns to normal a er about 100 min longer than the sensorial level regression from the S2 to S3 level ³⁵. It has been demonstrated that spontaneous urina on may be influenced by an intrathecal local anaesthe c; long-ac ng agents require a longer me to recover from urinary func on 36 .

Several surgical risk factors may increase the incidence of urinary reten on, such as anorectal surgery, inguinal hernia, orthopaedic [especially hip] surgery, abdominal surgery, instrumental delivery, prolonga on of labour and gynaecologic surgery ³⁷. Pa ent characteris cs showed that being predisposed to urinary reten on included the male gender, 50 years and older and the presence of urina on problems ³⁸. Besides neuraxial anaesthesia, some of the anaesthesia-related or intraopera ve factors are prolonga on of anaesthesia or surgery, increased intraopera ve fluid volume [>750mL], a required atropine, decreased body temperature and opioid-based anaesthesia, which may increase urinary reten on ^{38,39}.

Spinal anaesthesia may also contribute to this complica on by increasing or contribu ng to the requirements of at least several factors men oned above. Commonly used addi ves such as opioids or epinephrine may also increase the me leading up to urina ng. In a meta-analysis, hydrophilic opioids were more prone to contribute to urinary reten on than



lipophilic compounds, which are especially important for outpa ent surgery ⁴⁰. The gap between general anaesthesia and neuraxial blocks are decreased when systemic opioids are predominantly used for pain control [38]. The dura on of spinal anaesthesia performed with hyperbaric local anaesthe cs is shorter than more plain solu ons, which may also be preferred ⁴¹. Interes noly, when compared to the same intrathecal dose, more dilute solu ons of local anaesthe cs regressed earlier and regained bladder func on faster⁴². Short-ac ng local anaesthe cs, the administra on of which should be given in as low a dose as possible, a plain or hyperbaric solu on with no addi ves and avoiding an unnecessary increase of sensorial levels may decrease this complica on in suscep ble individuals within the outpa ent se ng. Indeed, a me culous review indicates that there is no risk involved in single shot spinal anaesthesia when such precau ons are taken⁴³. Bladder volume is alsoan important issue during admission to the intensive care unit. Single bladder catheteriza on may be necessary during the peri-opera ve period or immediately following surgery.

Ultrasound may precisely determine the bladder volume in adults and can be an important part of rou ne use in post-anaesthesia care units⁴⁴. A mul modal approach for postopera ve pain regarding the decrease of systemic opioids might be beneficial to avoid unnecessary hospital re-admissions. Non opioids, non-steroid an inflammatory drugs and other regional techniques such as wound infiltra on and peripheral nerve blocks have also been demonstrated to decrease urinary reten on⁴³.

Haematologic complica ons

Spinal haematoma following spinal anaesthesia is a severe complica on that requires early surgical interven on to prevent permanent neurological damage. Classically, the incidence of this condi on has been accepted as 1 in 220, 000 pa ents undergoing spinal anaesthesia, but the actual incidence remains unknown and is presumed to be on the increase. Advanced age, female gender, pa ents receiving drugs that influence coagula on, di culty in performing block and placement of the indvelling epidural catheter are men oned asrisk factors⁴⁵.

A study inves ga ng neurologic complica ons a er neuraxial block, performed in Sweden overa period of ten years, indicates an increased incidence in female pa ents undergoing hip fracture surgery – 1 in 22, 000 compared to 1 in 480, 000 when all pa ents were included⁴⁶. Haematoma was more frequently encountered with epidural anaesthesia

or catheter placement, because of the increased vascularity of the epidural space. The presence of haematomais frequently suspected in the case of an unexpected increase in the dura on of motor block or delay on recovery. Neurosurgery within eight hours a er the epidural haematoma is mandatory to regain motor func ons without neurologic harm. Should neurologic harm be suspected, imaging studies, including computed tomography or preferably MRI, should be implemented as early as possible. Unfortunately, neurologic outcomes have been poor for the majority of pa ents, even when surgery was performed within eight hours. Spinal catheters should be considered as epidural catheters, for which placement or removal requires strict adherence to withdrawal guidelines to avoid having an e ect on an coagula on therapy.

Many drugs interfere with blood clo ng, thereby requiring adherence to recommenda ons for the removal of neuraxial catheters A number of regional anaesthesia socie es have published their own recommenda ons^{47,48,49}. Recent evidence indicates spinal anaesthesia to be safe, provided the half-life of the drug or residual e ects are monitored. Nonsteroidal an - inflammatory drugs are considered safe, but concomitant drug use can increase the risk of haematoma. Therefore, pa ents receiving more than one drug a ec ng coagula on should be carefully evaluated. Addi onally, some herbal drugs only or in combina on with an coagulants can increase the risk of spinal haematoma⁴⁵.

Central neuraxial blocks in pa ents with pre-exis ng haematologic disorders or disease a ec ng coagula on do not appear to be a significant problem. In their review, *Choi and Brull*⁵⁰ inves gated the outcome of neuroaxial anaesthesia in pa ents with common bleeding disorders. A total of 78 spinal anaesthe cs, 53 diagnos c lumbar punctures and two combined spinal and epidural anaesthe cs were performed. No bleeding complica ons were observed, except in one infant, who was an unknown haemophilia A, developed spinal anaesthesia appears to be safe in pa ents with known bleeding disorders, provided that the status of coagula on is monitored. Although there is no



consensus concerning a safe platelet count, 50, 000 to 80, 000mm3 is generally considered a cri cal number for spinal or epidural anaesthesia. Individual pa ent assessment should be conducted in pa ents with lower platelet counts⁵¹. For more details, the reader is referred to the chapter on spinal haematoma included in this book.

Infec ouscomplica ons

Although bacterial meningi s following neuraxial anaesthesia is an uncommon complica on, in cases where it does occur it may result in severe harm, including permanent neurologic disability and death. The presence of a fever and neurologic disturbance may provide a di eren a on from PDPH. Epidural abscess is generally caused by skin flora; the bacteria most frequently involved is S. aureus. It is therefore prudent to ini ate treatment with synthe c penicillin, even in the absence of a posi ve culture. Other less common causes of infec on are aerobic and anaerobic streptococci, and anaerobic gram-nega ve bacilli. The incidence of meningi svaries between 1 in 50, 000 and mostly occurs as a result of airborne pathogens. The exact mechanism for how the microorganism reaches the spinalcord remains controversial. It may occur during prepara on or performing the block, witha droplet from medical personal is the predominant source. Infec on is more likely to occur in streptococci in most of the cases, emphasizing the need for strict adherence to precau ons while performing spinal anaesthesia⁵².

A case report and review from the literature indicates 179 cases of bacterial meningi s related to the central neuraxial puncture for any indica on covering the period 1952 to 2005, in which 54% was related to spinal anaesthesia and 5% was observed for the combined spinal and epidural technique, which included 15 obstetric pa ents. Technical di cul es during placement of the needle or repeated a empts to spinal anaesthesia appear to be contribu ngfactors⁵³.

Spinal anaesthesia in pa ents with coexis ng infec on is a controversial issue. In their study, *Gritsenko et al.* ⁵⁴ retrospec vely reviewed pa ents who had undergone removal of an infected prosthesis due to hip or knee arthroplasty performed under neuraxial anaesthesia to look for possible associa ons between periopera ve infec on and postopera ve neuraxial complica ons regarding meningi s or epidural abscess. Although higher incidence of posi ve joint culture or pus was found during these procedures, none of the pa ents included in the 474 cases demonstrated infec ous complica ons during the postopera ve period. A study performed by *Bader et al.* ⁵⁵ inves gated 319 obstetric pa ents with chorioamnioni s, eight of whom had bacteraemia, but none developed neuraxial infec on following neuraxial anaesthesia. A similar study of 517 pa ents with the same pathology, including 13 cases of systemic infec on, demonstrated no meningi s or epidural abscess [56].

These results indicate that the possibility of haematologic spread through the spinal cord from the remote site as an infec ous source is less likely to occur; regardless, clinicians are advised to perform the block under empirical an bacterial therapy.

Spinal anaesthesia in a pa ent with immunodeficiency is another instance where the presence of infec on has been observed to lead to posi ve CSF culture and infec on ⁵⁷. Therefore, performing spinal anaesthesia in such pa ents requires strict a en on and may be a empted in combina on with an bio c treatment. On the other hand, in a large prospec ve study including obstetric pa ents, general anaesthesia was associated with a higher incidence of surgical site infec on and postopera ve hospital stay compared to the use of spinal or epidural blocks⁵⁸.

Neurologic complica ons

A review by Brull et al.⁵⁹, which included a large series of neurologic complica ons, reported that the incidence of permanent neurological injury following spinal anaesthesia varied between O to 4. 2 per 10000 pa ents. In a French survey, permanent neurological injury other than that caused by haemorrhage was more common and included injury to the conus medullaris, and the es mated risk was calculated as 1:78 660 spinal anaesthesia pa ents; incidence was nearly half the amount in obstetric pa ents compared to the nonobstetricpopula on. Pre-exis ng spinal pathology or disease increases the incidence of postopera ve neurologic complica ons following neuraxial blockade. Repeated a empts or improper posi oning of pa ents may facilitate neurologic injury⁶⁰. Lumbar canal stenosis is another contribu ng factor for adverse neurologic outcome ⁶¹. The presence of scoliosis with or without prior surgery cons tutes di cul es for performing neuraxial anaesthesia. When compared to spinal blocks, the success rate was lower with epidural anaesthesia,



due totechnical di cul es and improper distribu on of local anaesthe cs⁶². Although the precise mechanism was not determined, hydrosta c pressure performed during an epidural block was indicated as a possible source of injury. On the other hand, direct needle trauma appears to be one of the preventable reasons for neurological complica ons. It is best to withdraw the needle in the case of paraesthesia, which is highly associated with postopera ve radiculopathy and repea ng local anaesthe c injec on should be avoided in order to prevent toxic concentra ons in the spinal cord ⁶³.

Reynolds⁶⁴ reported a series of cases of conus medullaris injury that including one nonobstetric and six obstetric female pa ents, resul ng in long-las ng neurologic damage. Spinal anaesthesia was performed in three of the pa ents, while combined spinal and epidural anaesthesia was accomplished in the rest of the other pa ents. Only one pa ent su ered pain during needle placement. The possible reasons for this were indicated as misplacement of the needle at the lower end of the spinal cord, misiden fica on ofTu er's line or that the arachnoid membrane may have been a ached to the conuslike a web. Author concluded that Tu er's line was an unreliable method for iden fying the correct intervertebral level⁶⁴.

Possibly, the addi on of cutaneous and subcutaneous ssue over the crista iliaca in obese pa ents, or in the case of pregnancy, may erroneously lead to performing a higher intervertebral space for needle placement. Indeed, Broadbent et al. 65 demonstrated that an anaesthesiologist incorrectly iden fied the correct intervertebral space by palpa on. Assuming the correct intervertebral space was correct in only 30% of pa ents using palpa on and in 71% using ultrasound [66]. In a MRI study of 690 pa ents, Kim et al. 67 indicated that cau on should be exercised when selec ng the appropriate intervertebral space, especially in obese and elderly pa ents. The level of conus medullaris might be lower than expected in female pa ents with thoracic vertebral compression fractures ⁶⁸. These points have also been highlighted along with determining lateral needle devia on or placement as a source of possible reasons for neurological injury in a study group by the American Society of Regional Anesthesia [ASRA]; clinicians are advised to especially be aware of challenging Surface anatomical changes ⁶⁹. Ultrasound imaging can also be used to guide proper

accomplishment of neuraxial anaesthesia. Although promising results have been published on facilita ng neuroaxial anaesthesia in di cult cases by means of decreasing the meand number of a empts⁷⁰, to date, it is not yet possible to conclude that using an ultrasound guide may decrease complica ons⁶⁹. Anaesthesiologists should also be cau ous concerning pa ents with preexis ng comorbidi es, such as peripheral vascular disease and diabetes mellitus, which may present subclinical neuropathies that could predispose the pa ent to neurological deficits following spinal anaesthesia⁷¹.

Skin an sep cs like chlorhexidine have proven to be superior to iodopovidone-based solu ons.ASRA advises using chlorhexidine in an alcohol solu on prior to all regional anaesthe cinterven ons to prevent infec ous complica ons ⁷². In a recent retrospec ve study, the neurological complica on rate when using chlorhexidine was found to be similar to the findings of other surveys⁵⁷. However, using chlorhexidine is not en rely devoid of risk. In an editorial, Bogod 73 published two cases of chlorhexidine inducing permanent neurological injury. In one case, chlorhexidine solu on was inadvertently administered into the epidural space. In the second case, 0.1 mLan cep cofsolu on (chlorhexidine) mixed with a local anaesthe c was wrongly administered to the subarachnoid space. The author advised using a spray formula on for skin prepara on, warning against high concentra ons [more than 2%]. Applying one pu was concluded to be su cient and emphasize was placed on wai ng for the skin to dry. Figure 2 shows medullaris cone injury secondary to a empted spinal anaesthesia in an obstetric pa ent undergoing a caesarean sec on. This pa enthad severe pain during bupivacaine injec on. The injury was managed with steroids. Final neurological damage was minimal.

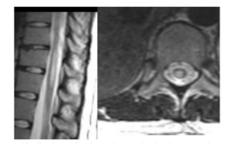


Figure 2 Sagi al and axial images of the conusmedullaris T2 weighted FSE in which a high signal is noted at the centre posi on, with conusoedema and/orhaemorrhagecausedby a spinalneedle (Source: www anestesia-dolor.org).



Neurological diseases

Pa ents with pre-exis ng neurological diseases such as mul ple sclerosis, amyotrophic lateral sclerosis, or a post-poliomyeli s condi on have previously been considered as rela ve contraindica ons for neuraxial anaesthesia. A double crush phenomenon was described to explain the deteriora on of neurological disease in the case of vulnerable neurons. It is believed that mechanical trauma caused by a needle or catheter, toxicity induced by local anaesthe cs or neural ischaemia due to addi ves could worsen the pa ent's neurological status. Increased stress may induce inflamma on deteriorates the dinical course, that may confuse with neural injury due to the procedure. However, recent evidence has demonstrated that spinal anaesthesia might be an op on in thispa entgroup⁷².

Neuraxial anaesthesia in pa ents with spinal canal pathology, including lumbar disk disease, spinal stenosis or previous surgery, is another issue that requires special concern. In a retrospec ve review, *Hebl et al.*⁷⁴ found that pa ents with pre-exis ng spinal canal pathology demonstrated a higher rate of neurological complica ons. However, a lack of control groups receiving general anaesthesia presents a di culty for describing results pertaining to whether complica ons occur due to surgery or because of the natural progression of disease. Addi onally, due to data combina ons, it is also impossible to conclude the par cular role of single shot spinal anaesthesia. Trauma and complica ons may be more common when using large gauge Tuohy needles, or during catheter placement. Epidural anaesthesia and catheter placement in pa ents with previous spinal surgery appear to be more complicated, even when performed by experienced hands ⁷⁵. These pa ents require special a en on in terms of evalua ng the use of neuraxial anaesthesia, preopera ve neurological evalua on and special care in order to prevent addi onal injuries.

In rare instances, silent pathologies involving the spinal column may induce acute postopera ve neurological complica ons such as tuberculosis ⁷⁶ or unrecognized spinal tumour ⁷⁷. Pa ents with coexis ng or previous low back pain and paraesthesia or neurologic deficits should be carefully assessed preopera vely and their evalua on should indude a detailed neurologic examina on and radiological images. Moreover, pa ents with neural tube defects should be assessed earlier to determine the conus medullaris level or other possible associated

anomalies to decide whether neuraxial techniques will provide them with safe anaesthe cop ons.

Pa ents like those described above are at higher risk of neurological complica ons than the rest of the general popula on⁷⁸. Postopera ve neurological complica ons were observed in a pa ent presen ng adhesive arachnoidi s, extensive syringomyelia and agiant arachnoid cyst in the pa ent had been managed with a combina on of spinal and epidural anaesthesia. These rare complica ons were linked to a reac on caused by the subarachnoid, the epidural drug, or as a result of catheter induced inflamma on or trauma⁷⁹.

The images in Figure 3 show a case of mul ple neurofibromatosis with intrathecal par cipa on, which was managed with uncomplicated spinal anaesthesia.

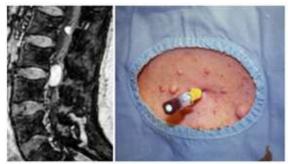


Figure 3. Forty year old pa ent with mul ple neurofibromatosis. She was anesthe zed successfully with spinal anaesthesia for an abdominal hysterectomy. The back of the pa ent showsnumerouskin tumours and some cafeaulait spots. The sagi al image of the lumbar spine in post-spin echo fat-suppressed contrast demonstrates intradural solid tumours in cauda equina nerves (Source: www.anestesia-dolor. org).

Cardiacarrest and periopera ve death

Bradycardia and cardiac arrest are the most worrisome complica ons related to spinal anaesthesia. The incidence of these condi ons has been observed to be higher with spinal block in comparison with general anaesthesia. Pa ents are generally healthy, ASA class I or II, athle cand male with parasympathe covertones.

The influence of cardio-accelerator fibres origina ng between T1 to T4 plays a crucial role in maintaining blood pressure and heart rate according to the level of anaesthesia induced by spinal block, depleted vascular volume or insu cient replacement with fluids, and the presence of deep seda on is considered a risk factors for bradycardia and cardiac arrest. Surgical interven on may also trigger bradicardia and cardiac arrest by vagal discharge or emboliza on. In the case of severe bradycardia, early administra on of epinephrine is important, especially in unresponsive cases to atropine



and ephedrine that should be administered previously [80]. Most pa ents are monitored in the opera ng theatre, therefore early recogni on of bradicardia and cardiacarrest and interven on ispossible. Survival rate is higher in pa ents with cardiac arrest observed during spinal anaesthesia compared to cases using general anaesthesia.⁸¹.

A study by *Chatzmichali* and colleagues ⁸² showed that assessment of heart rate variability in the preopera ve period may help to determine periopera ve severe bradycardia. Clinicians must be cau ous when performing deep intravenous seda on, especially in pa ents with increased body weight, since it may lead to death in the early postopera ve period.

Miscellaneous complica ons

Myodonus occurs rarely as a complica on of spinal anaesthesia in the postopera ve period. It may commonly be observed in the presence of systemic illness, drug use or with a preexis ng vitamin B deficiency. Although the underlying mechanism for myodonus in this instance is unclear, the possibility of subclinical neuropathy has been men oned ⁸³. Longterm treatment, including neurolep cs and benzodiazepines, might be required to relieve the symptoms of myodonus ⁸³.

Exposure to the halogenated compounds during general anaesthesia is an interes ng topic, especially in the context of the brain, which is currently under development as it relates to aged pa ents. Spinal anaesthesia appears to be safe for preven ng postopera ve delirium andcogni ve dysfunc on in the elderly, especially when addi onal measurements such as early pain management, supplemental oxygen, fluid, caloricreplacement and morphine avoidance are applied ⁸⁴.

Table 1 is a summary of the complica ons of neuraxial anaesthesia found in Finland. The authors reported 1:17 741 spinal block cases and 1:24 285 cases of epidural blocks. Complica ons were more severe in the applica on of spinal anaesthesia.

Table 1. Severe complica ons associated with epidural and spinal anaesthesia $^{\rm 85}$			
Claim mo ve	Spinal	Epidural	Total
Cardiacarrest	2[2]	<u>0</u>	<u>2</u>
Neurological	<u>31 [19]</u>	<u>7 [4]</u>	<u>38</u>
_Infec_ous	<u>4 [4]</u>	<u>6[2]</u>	<u>10</u>
Local anaesthe cs acute toxicity	Ο	2[2]	2
Opioid overdose	0	1 [1]	1
	9	8	17
Others	T3	3	16
Pa ents who died contracted permanent neuronal damage ————————————————————————————————————			

The authors reported 1:17 741 spinal block cases and 1:24 285 cases of epidural blocks. Complica ons were more severe in the applica on of spinal anaesthesia. Is a summary of the complica ons of neuraxial anaesthesia found in Finland.

Conclusions

Various complica ons may occur during spinal anaesthesia and are widely related to the procedure itself or drugs used during the procedure. These complica ons occur with di ering incidence and in the case of at least some, appear to be inevitable and to be expected due to the invasive nature of the blockade. Many of these complica ons can be reduced with me culous a en on to the details during the performance of the spinal block. The procedure may be rendered more pa ent-oriented and convenient by selec ng the appropriate technique, drugs and their doses. It is of prime importance that the incidence of hypotension should be decreased, as this can induce serious adverse outcomes. Ultrasound guides may be helpful for decreasing complica ons in di cult cases. In order to decrease serious complica ons, pa ent selec on and adherence to the guidelines appears to be fundamental. (such as pa ents with previous lumbar surgeryasmen oned).

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