



# Lumbar puncture Mastery Pre-learning materials



## Authors:

Callum Mutch Karen Macsween Ian Morrison Andrew Page Rustam Al-Shahi Salman James Tiernan

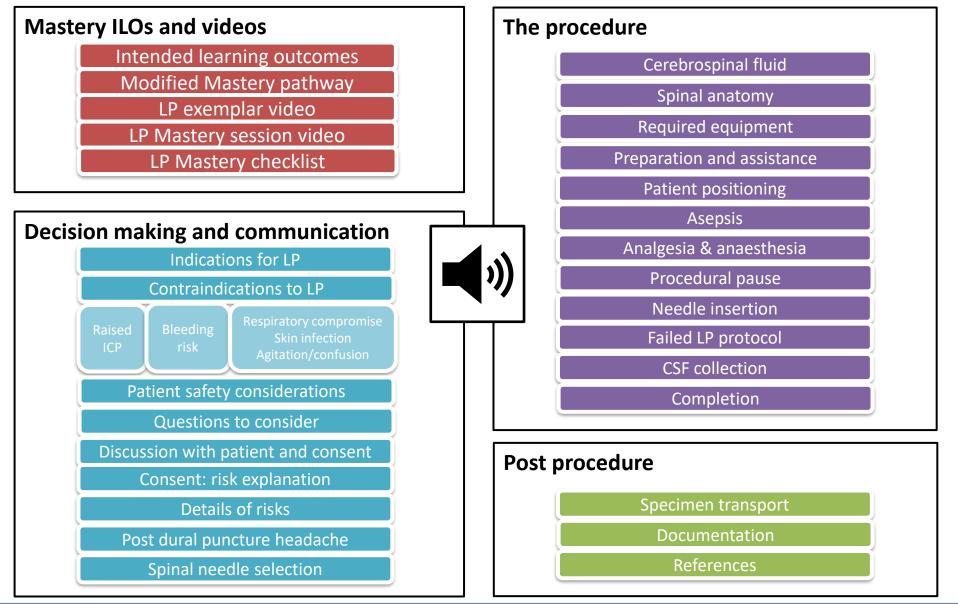
Autumn 2023

Medical Education Fellow Consultant Medical Microbiologist Simulation Technician Consultant haematologist Consultant neurologist NHS Lothian Mastery Lead



*Tip* Clicking the Home icon at the bottom right of each page will return you to this slide at any time.





Click on any button to visit the topic or click next to continue.



By the end of this Mastery pre-learning package you will be able to:

- 1. Describe the indications for performing a lumbar puncture and its use in clinical practice
- 2. Explain the risk assessment, patient safety concerns and contraindications to lumbar puncture
- 3. Describe the potential complications of the procedure and the basic principles of their management.
- 4. Explain the practicalities of performing a lumbar puncture in a safe and structured fashion.
- 5. Recognise your limitations and when to obtain help from a senior clinician.
- 6. Describe the process for ensuring the correct CSF results are requested and obtained.



Mastery learning pathway

))

# **Modified Mastery Pathway**

**Enhanced Pre-Learning** 

**Peer-Assisted Deliberate Practice** Mastery **Facilitated Furthering** Session Simulated Performance & Assessment



# Lumbar puncture

Please watch the below LP exemplar video [20 minutes]



Lumbar Puncture (scot.nhs.uk)



# Lumbar puncture

Please watch the below LP learner Mastery session video [49 minutes]



Lumbar Puncture (scot.nhs.uk)



#### Lumbar puncture checklist

This is the checklist you will be assessed against at your Mastery session

Consider these steps as you read through the rest of the prelearning

Phase 1 – Preparation + Positioning
Identifies correct patient and lists contra-indications (Sheet 1, Q1)
List core consent topics (Sheet 1, Q2)
Lists essential equipment + ensures trained assistant present (Sheet 1, Q3)
Describes optimal position (left lateral)
Optimises / Describes ergonomics (seat/bed height/equipment)
Identifies and marks insertion site (L3/4 or L4/5)
Phase 2 – Asepsis + Anaesthesia
Puts on surgical Mask + Hat / Washes Hands / Gown + Sterile Gloves
Applies antiseptic skin wash + allows to dry (non-touch)
Avoids contamination of equipment and gloves by the cleaning solution
Drapes the patient (non- touch +/- tape by assistant)
Infiltrates local anaesthetic – skin bleb +/- deeper
Phase 3 – Procedural Pause
Performs final equipment check + assembles manometer
Performs 3-point check: Patient, Assistant and Clinician
Phase 4 – Insertion
Inserts introducer
Inserts pencil-point spinal needle
Obtains CSF flow within 3 attempts
Measures opening pressure
Collects CSF: 3 samples of at least 2mls (utilises assistant)
Phase 5 – Anchoring + Dressing
Not applicable
Phase 6 – Completion + Global Points
Describes aftercare: Samples / Analgesia / Pt instructions / Documentation
Maintains control of introducer and needle throughout
Demonstrates safe sharps management throughout
Maintains asepsis throughout
Demonstrates effective communication with assistant throughout



DIAGNOSIS OF LIFE-THREATENING CONDITIONS

Subarachnoid haemorrhage

# Acute Central Nervous System (CNS) infections

- meningitis
- encephalitis

### DIAGNOSIS OF NEUROLOGICAL CONDITIONS

Other meningitides e.g. tuberculosis, carcinomatous

Demyelination/ Inflammatory conditions e.g. Guillain Barre Syndrome, Multiple Sclerosis, Vasculitis

Raised intracranial pressure e.g. Idiopathic intracranial hypertension, cerebral venous sinus thrombosis

Cognitive decline e.g. Creutzfeldt-Jakob Disease



All potential contraindications should be assessed and a decision made about the risk versus benefit of the procedure for each individual.



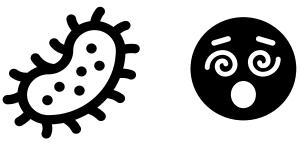
Raised ICP



**Bleeding Risk** 



Respiratory Compromise



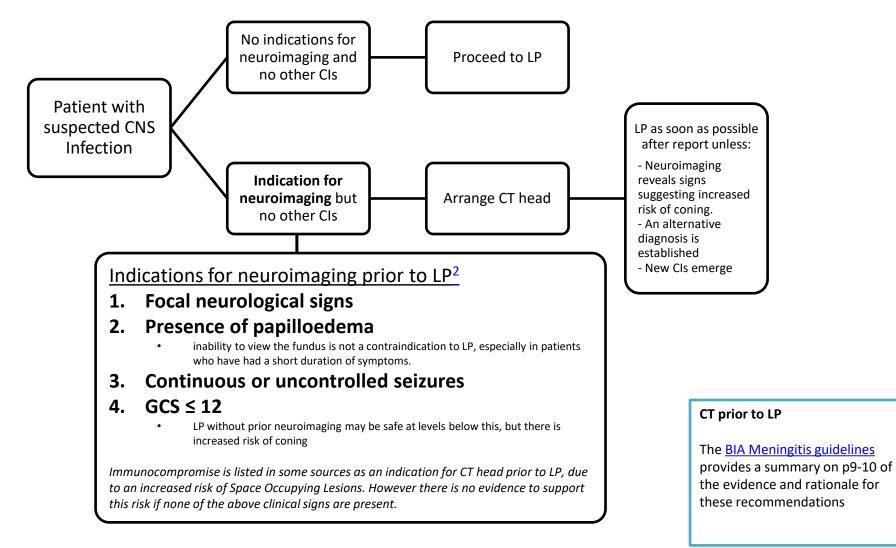
Skin infection at site

Agitated/ confused patient

More details to help you assess these are on the following slides



# **MENINGOENCEPHALITIS**





# **MENINGOENCEPHALITIS**

The gold standard of care for suspected CNS infection would include:

- 1. Urgent clinical assessment
  - The need for a rapid LP has to be weighed against the desire to start antimicrobial treatment urgently<sup>1</sup>.
- 2. Rapid lumbar puncture prior to antimicrobials
  - Even if antimicrobial treatment has been initiated, a LP should still be performed as soon as possible, preferably within 4 h of commencing antibiotics, to help identify the cause of meningitis<sup>2</sup>.
  - The culture rate (and PCR positivity) can drop off rapidly after that time and it can become difficult to identify the causative bacteria in cases of bacterial meningitis<sup>1</sup>.
- 3. Prompt antimicrobials
  - Intravenous antibiotics should be given promptly in hospital as there is evidence that delays increase mortality<sup>3, 4, 5</sup>.



#### **OTHER INDICATIONS**

In suspected SAH CT head should be performed prior to LP. LP should be delayed until >12 post headache onset to improve diagnostic yield



# Lumbar Puncture and Bleeding Risk

A brief guide to managing antiplatelets and anticoagulation in patients requiring lumbar puncture (LP).

## Patient requires lumbar puncture

### **Consider Bleeding Risk**

Performing an LP with coagulopathy increases the risk of spinal haematoma

#### **Consider Thrombosis Risk**

Suspending antithrombotic treatment comes with an increased risk of thrombosis

The following slides are not a local protocol and in cases of uncertainty advice should be sought from the local Haematology team

*Reference<sup>6</sup>: Dodd, K. C. et al. (2018) 'Periprocedural antithrombotic management for lumbar puncture: Association of British Neurologists clinical guideline', Practical Neurology, 18(6), pp. 436–446. doi: 10.1136/practneurol-2017-001820.* 



# Lumbar Puncture and Bleeding Risk

A brief guide to managing antiplatelets and anticoagulation in patients requiring lumbar puncture (LP).

# Patient requires lumbar puncture

### **Consider Bleeding Risk**

Performing an LP with coagulopathy increases the risk of spinal haematoma

- Routine coagulopathy testing in unselected patients is not recommended
- Consider a patient's risk of haemorrhage on an individual basis
- Consider checking platelet count is >40 x10° and PT/APTT within normal range if high risk for coagulopathy:
  - liver or renal failure
  - heparin treatment for >5 days
  - haematological disorder
  - disseminated intravascular coagulation
  - personal/family history of unexplained bleeding

If at high risk for bleeding discuss with haematology team or consider postponing LP

### **Consider Thrombosis Risk**

Suspending antithrombotic treatment comes with an increased risk of thrombosis

#### Haematology comments:

Almost all patients will have a FBC +/- coagulation screen performed prior to LP, so always check the results of these if available.

Although a coagulation screen is not required routinely, it should be performed in the context of thrombocytopaenia as the most likely cause of this in the context of CNS infection would be consumption/DIC

In the context of platelets  $<40 \times 10^9$  consideration of platelet transfusion

Ultimately your decision is balancing the risk of delaying an LP to perform a coagulation screen versus the risk of missing a coagulopathy and increased risk of spinal haematoma



.

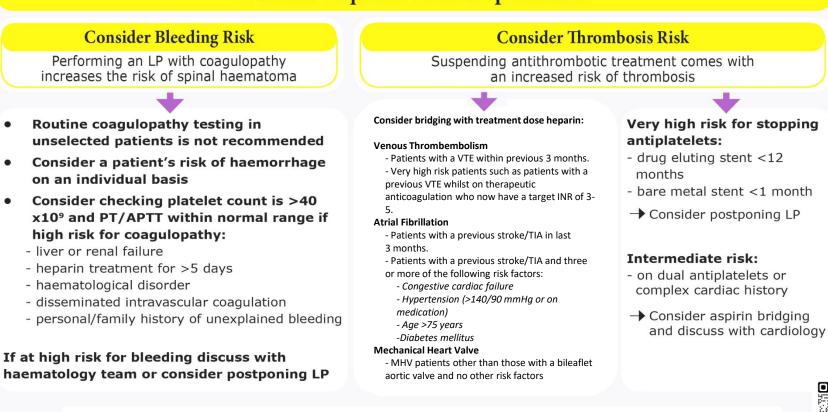
•

.

# Lumbar Puncture and Bleeding Risk

A brief guide to managing antiplatelets and anticoagulation in patients requiring lumbar puncture (LP).

# Patient requires lumbar puncture





Bridging guidelines here: Keeling, D., Tait, R.C., Watson, H. and (2016), Peri-operative management of anticoagulation and antiplatelet therapy. Br J Haematol, 175: 602-613. https://doi.org/10.1111/bih.14344

.

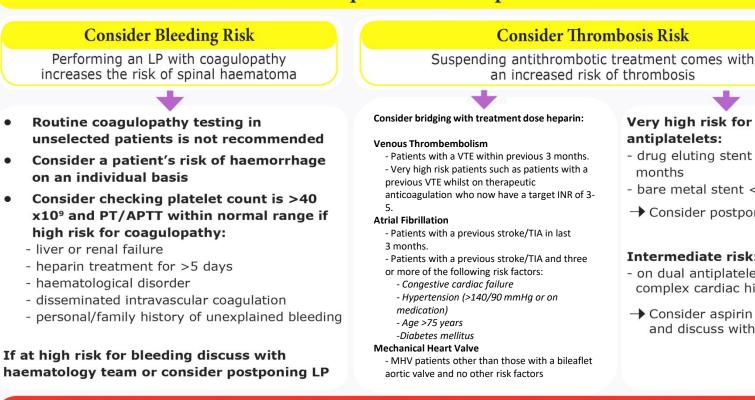
•

.

# Lumbar Puncture and Bleeding Risk

A brief guide to managing antiplatelets and anticoagulation in patients requiring lumbar puncture (LP).

# Patient requires lumbar puncture



Urgent LPs: Patients may require an urgent LP outside of these time frames. Discuss with haematology about reversing warfarin and some DOACs. In other situations it may be decided that the benefit of an LP outweighs the increased bleeding risk, but the patient must be informed of this risk and be carefully monitored for new neurological symptoms or signs.

#### Very high risk for stopping antiplatelets:

- drug eluting stent < 12
- bare metal stent <1 month
- → Consider postponing LP

#### Intermediate risk:

- on dual antiplatelets or complex cardiac history
- → Consider aspirin bridging and discuss with cardiology

## **Bleeding risk: discontinuing antiplatelets**



Discontinuing medications in patients with normal renal function

<u>Haematology comments:</u> Consider aspirin cover in patients at high risk of thrombosis who were on alternate antiplatelets

All patients on antiplatelets should have a FBC checked to assess for thrombocytopaenia prior to LP

*This slide is an amalgamation of the following guidelines:* 

Harrop-Griffiths, W. et al,. (2013), Regional anaesthesia and patients with abnormalities of coagulation. Anaesthesia, 68: 966-972. doi.org/10.1111/anae.12359

Dodd, K. C. et al. (2018) 'Periprocedural antithrombotic management for lumbar puncture: Association of British Neurologists clinical guideline', Practical Neurology, 18(6), pp. 436–446. doi: 10.1136/practneurol-2017-001820.



# Bleeding risk: discontinuing anticoagulants

### Discontinuing medications in patients with normal renal function

#### Haematology comments:

Any patient whose anticoagulation is being stopped is likely to need thromboprophylaxis unless contraindicated

When stopping warfarin, prophylaxis should be started once the INR is <2, not when the first dose is missed.

*If LP is essential in the context of Fondaparinux treatment then this can be discussed with haematology* 

This slide is an amalgamation of these guidelines:

Harrop-Griffiths, W. et al,. (2013), Regional anaesthesia and patients with abnormalities of coagulation. Anaesthesia, 68: 966-972. doi.org/10.1111/anae.12359

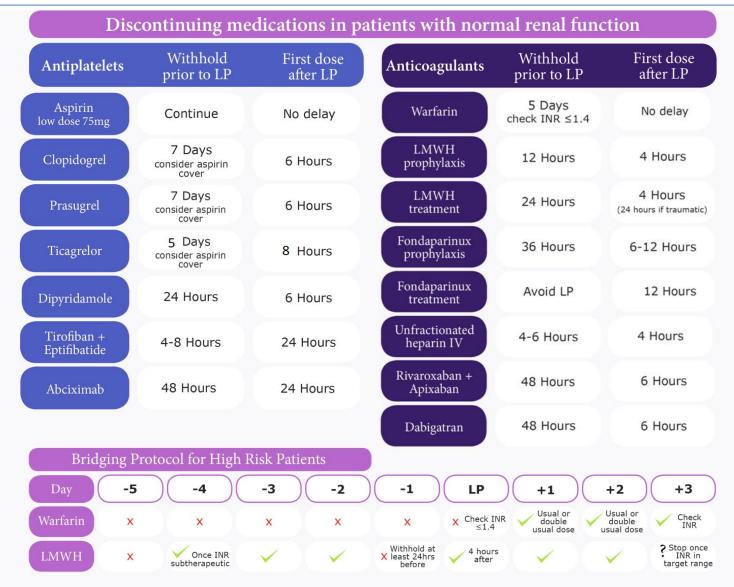
Saja, K. Addendum to the guideline on the peri-operative management of anti-coagulation and anti-platelet therapy. Br J Haematol. 2022; 197: 188– 189. https://doi.org/10.1111/bjh.18114

Dodd, K. C. et al. (2018) 'Periprocedural antithrombotic management for lumbar puncture: Association of British Neurologists clinical guideline', Practical Neurology, 18(6), pp. 436–446. doi: 10.1136/practneurol-2017-001820.

Anticoagulants	Withhold prior to LP	First dose after LP
Warfarin	5 Days check INR ≤1.4	No delay
LMWH prophylaxis	12 Hours	4 Hours
LMWH treatment	24 Hours	4 Hours (24 hours if traumatic)
Fondaparinux prophylaxis	36 Hours	6-12 Hours
Fondaparinux treatment	Avoid LP	12 Hours
Unfractionated heparin IV	4-6 Hours	4 Hours
Rivaroxaban + Apixaban	48 Hours	6 Hours
Dabigatran	48 Hours	6 Hours



# **Bleeding risk: bridging**

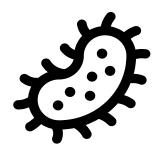


▲ 18



## **RESPIRATORY COMPROMISE**

- Patients with any form of respiratory compromise (especially respiratory muscle weakness) present significant risk in the context of invasive procedures.
- Particular attention must be paid to positioning, ventilatory and physical support of such patients.
- If the patient is deemed fit enough, by senior clinicians, to undergo such a procedure, it should ideally be performed in the upright position.



## **SKIN INFECTION**

- Skin infection, cellulitis or suspected epidural abscess at the site of the LP can potentially introduce infection into the CSF space.
- If any concerns, delay procedure and seek help.



## AGITATED OR CONFUSED PATIENT

- There is a higher chance of failure, trauma and infection if the patient is unable to remain still.
- A senior, experienced clinician should attempt the LP for an agitated patient.
- The patient may require mild sedation and discussion with anaesthetists.



### Click any of the icons to return and review that contraindication



**Raised ICP** 



**Bleeding Risk** 



Respiratory Compromise





Skin infection at site

Agitated/ confused patient



When performing lumbar puncture, the goal is to safely obtain diagnostic information having removed or minimised any potential risk factors.

If there is any concern that significant risk of harm may compromise patient safety, delay the procedure and seek senior advice.



# **Resuscitation equipment**

- Resus trolley available
- Consider IV access preprocedure



# Competent practitioner

• Supervisor available for practitioner, if required



# Skilled assistant

- Competent in equipment checking and ensuring patient comfort
- Familiar with environment and procedure



*Consider these questions prior to any invasive procedure:* 

Does it need to be	Does it need to be	Am I competent to
done?	done now?	do this?
Is supervision/ assistance available?	Am I familiar with the equipment?	Does the patient have capacity to consent to the procedure?



The patient should be made aware:

- Why they are having the procedure
- The benefits
- <u>The potential risks/ complications</u> (outlined on next 2 slides)
- The alternatives to the procedure.

Once this has been done, informed consent should be sought and documented. There is a consent form available in NHS Lothian which can be used for written consent for lumbar puncture.

If the patient does not have capacity to give informed consent, and the procedure is deemed clinically necessary, ensure an Adults with Incapacity form is completed.

Always establish whether the patient has any known allergies prior to the procedure - they may have an allergic reaction to local anaesthetic or antiseptic skin preparation.



# What are the risks?

## A recommended script for discussion with patients:

The most common side effects of having a lumbar puncture are<sup>Z</sup>:

- Short-lived back pain (1 in 6 people)
- Shooting leg pain during the procedure (1 in 9 people)
- Headache (1 in 10 people).

Sometimes the first attempt at the procedure is not successful. In this case we may need to repeat it.

There are also some more serious, rare risks<sup>8</sup>:

- Infection: Abscess (1 in 47,000 people) and Meningitis (less than 1 in 200,000 people)
- Serious bleeding: Spinal haematoma (less than 1 in 100,000 people)
- Permanent nerve damage (1 in 100,000 people).

Note that the best evidence is derived from National Audit of spinal procedures involving mainly spinal anaesthesia which generally use large bore needles and consequently have higher risks of bleeding and infection. The risks are likely lower for diagnostic lumbar puncture<sup>8</sup>



# Risks from Lumbar puncture and management

Complication	Estimated incidence	Clinical presentation	Recommended action
Post dural puncture headache	1 in 10	Headache following lumbar puncture which is usually worse sitting/standing and improved on lying flat.	See <u>next slide</u>
Backache <sup>Z</sup>	1 in 6	Mild localised or generalised lumbar pain	Simple analgesia, reassurance and worsening advice for symptoms of more severe complications below
Infection: <b>local cellulitis</b> <sup>Z</sup>	No frequency data	Spreading cutaneous erythema without evidence CNS infection	IV antimicrobials as per local guidelines Consider deeper infection
Infection: Vertebral canal abscess & Discitis/Osteomyelitis <sup>8</sup>	1 in 47,000	Local signs of infection +/- systemic signs of infection. Delayed presentation of weeks/months possible	MRI spine Blood cultures Infectious diseases, neurosurgery and neuroradiology discussion
Infection: Meningitis/ meningoencephalitis <sup>8</sup>	<1 in 200,000	Pyrexia, neck stiffness, confusion, altered consciousness. May be atypical presentation.	LP Infectious diseases referral
Spinal or extradural haematoma <sup><u>8</u></sup>	1 in 117,000	Persistent/severe back pain Lower limb neurological deficit	Urgent MRI spine Neurosurgical discussion
Cerebral herniation	Risk theoretical based on historical studies <sup>9</sup>	Reduced GCS, Coma, dilated fixed pupils, Bradycardia and hypertension (Cushing's reflex)	ICU referral Neurosurgical discussion



#### Definition and aetiology

Headache following a lumbar puncture (LP) typically has a low-pressure phenotype; i.e. worse upright and better lying flat. It is usually caused by a dural tear sustained at the time of LP and does not relate to the volume of cerebrospinal fluid (CSF) taken. In most cases it is self-limiting although a few patients may require a blood patch for persistent headache, and rarely the low pressure may be associated with the development of subdural haematomas.

# Practices associated with reduced risk of post-LP headache



Finer gauge needles <sup>10, 11</sup>	Decreased risk with smaller gauge needles. Although smaller needles take longer to collect the required samples, this is not associated with patient discomfort <sup>24</sup> 22G is likely the ideal size, balancing collection duration with reduction in risk
Atraumatic needles <sup>11, 12</sup>	Atraumatic (e.g. Whitacre/Sproute) needles should be used first line. There is high quality evidence that these lead to a lower risk of post-LP headache than traumatic (e.g. Quincke) needles
Orientation of the bevel of the needle <sup>13, 14</sup>	Use of needle in a transverse plane (perpendicular to the longitudinal axis) This is less relevant when using an atraumatic needle
Replacement of the stylet 15	Unlikely to be relevant with atraumatic needles.
Experience and number of attempts <sup>16</sup>	Fewer attempts at dural puncture is associated with a decreased incidence of headache after lumbar puncture



#### Definition and aetiology

Headache following a lumbar puncture (LP) typically has a low-pressure phenotype; i.e. worse upright and better lying flat. It is usually caused by a dural tear sustained at the time of LP and does not relate to the volume of cerebrospinal fluid (CSF) taken. In most cases it is self-limiting although a few patients may require a blood patch for persistent headache, and rarely the low pressure may be associated with the development of subdural haematomas.

# Practices NOT proven to reduce risk of post LP headache



Reducing the volume of CSF taken <sup>17</sup>	There is no evidence that the amount of CSF removed influences the incidence of post LP headache
Bed rest <sup>18, 19</sup>	Patients are often advised to lie recumbent for a period of time after an LP but there is no evidence that this reduces the risk of headache.
Hydration <sup>19</sup>	There has only been one study looking at fluid post LP as a preventative strategy and it showed no difference between those who took 1.5 L and those who had 3 L post LP.
Caffeine	IV caffeine can be used to treat post LP headache but there is no evidence that either oral or IV caffeine can prevent the headache. Oral caffeine can be recommended if the patient develops a headache, but this is not evidence based.



## **Needle selection**

Lumb Lumb	bar puncture any rentional		or	Atraum needle Lumbar p with any a (pencil po needle	uncture atraumatic		
	Convent	tional needl	е	Atraur	natic needle		
	Strong	Weak		Weak	Stron	g	
We recommend the use of atraumatic over conventional needles							
Comparison of benefits and harms							
Source <sup>20</sup> : <u>Atraumatic</u>	Favours convention	nal needle 🧹	No important diffe	erence	Favours atraum	atic needle	
(pencil-point) versus			Events per 1000 p	people		Evidence q	uality
conventional	Postdural puncture headache	98	59 f	ewer	39	★★★★ Hi	
needles for	Need for epidural blood patch	24		ewer	12	★★★★ Hi	
lumbar puncture:	Backache	166	No important diffe		159	★★★★ Hi	
a clinical practice	Hearing disturbance	53		ewer	13	★★★★ Hi	-
guideline   The	Nerve root irritation	126		ewer	89	★★★★ M	
<u>BMJ</u>	Hospital for fluids or analgesia	39		ewer	17	★★★★ Hi	
	Failed lumbar puncture	38	No important diffe	erence	33	<b>★★★★</b> Hi	gh



# **CEREBROSPINAL FLUID (CSF)**

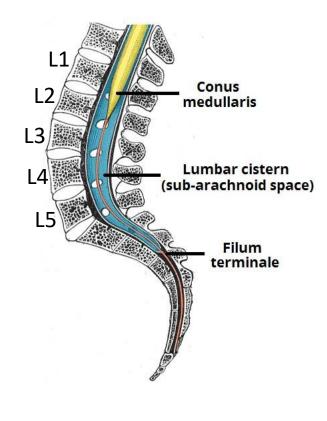
- provides hydromechanical protection of the central nervous system and plays a prominent role in brain development and regulation of brain interstitial fluid homeostasis, influencing neuronal functioning.
- The mean CSF volume in adults is 150ml, with 25ml in the ventricles and 125ml in subarachnoid spaces.
- Approximately 500-600mls of CSF is produced by the choroid plexus (and other cells) over 24 hours i.e. the entire volume is renewed about four times every day.
- Approximately 15-17ml of CSF can be safely taken in adults (~250-280 drops)
- The normal range for adult CSF opening pressure (measured with the patient in a lateral position) is 10-25cmCSF.

	Mean CSF production rate (ml/h)	CSF volume (mls)	Safe CSF volume to take at LP (mls)
Adult	22	150-170	15–17
Adolescent	18	120-170	12-17
Young child	12	100-150	10-15
Infant	10	60—90	6—9
Term Neonate	1	20—40	2—4

Table: Estimates of CSF production rate and safe volume to take<sup>27</sup>

### The spinal cord in adults

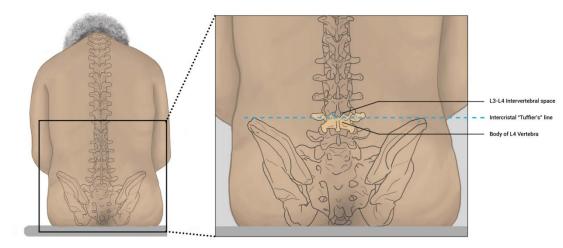
Image source (used with permission of creator) and to learn more: <u>The Spinal Cord - Meninges -</u> <u>Vasculature - TeachMeAnatomy</u>

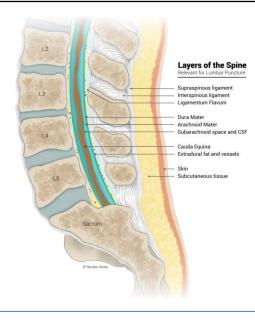




## SURFACE ANATOMY

An imaginary line connecting iliac crests at the level of the L4 vertebral body in most patients. Sometimes called Tuffier's line.





## **INTERNAL ANATOMY**

Sagittal section. The spinal needle will pass through skin, supraspinous ligament, interspinous ligament, ligamentum flavum and into the epidural space

Click the buttons below to view a 3D model:

Images and models © Renske Hoste // www.renskehoste.com

Animated needle Annotated model



# The procedure: required equipment & resources 1

The following should be available before commencing the procedure.



#### **PPE & Arranging equipment**

Full asepsis should be maintained throughout the procedure.

Use of surgical gown, mask, hat is recommended. There is evidence of iatrogenic infective complications following spinal procedures<sup>8</sup>

#### **ARRANGE EQUIPMENT**

- Put on head-cap and face-mask prior to opening sterile equipment.
  - In the rare cases of epidural abscess most reports identify bacteria that are found to have come from the operator's oral flora.<sup>8</sup>
  - Oral flora can contaminate samples if a mask is not worn
- Open sterile pack onto procedural trolley
- Open procedural equipment onto the trolley
- Ensure trolley on correct side for clinician

#### Skin cleaning<sup>21</sup>

Should be performed with 0.5% Chlorhexidine + 70% ethanol (Hydrex clear)

- Iodine has less antiseptic effect
- 2% Chlorhexidine has been associated with arachnoiditis
- Avoid contaminating the spinal needle with chlorhexidine solution

21. Wash & go – but with what? Skin antiseptic solutions for central neuraxial block - Checketts -2012 - Anaesthesia - Wiley Online Library

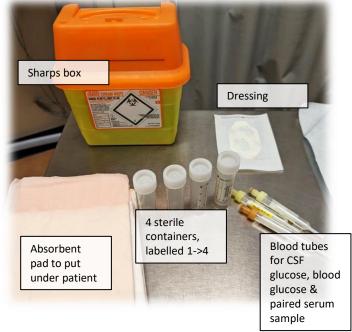




# The procedure: required equipment & resources 2

The following should be available before commencing the procedure.

Dressing pack: sterile gauze, pot for chlorhexidine Sterile drape 5ml syringe for lidocaine Applicators for Needles for chlorhexidine drawing up Atraumatic Sterile lidocaine, skin needle, 22G bleb and manometer or 25G deeper infiltration



Not pictured:

- 1. Experienced And Skilled Assistant
- 2. Stool/Chair For Operator To Sit On During Procedure
- 3. Pillows (X2)
- 4. Procedural Trolley
- 5. Lidocaine 1% Or 2% 5ml Ampoule
- 6. Black Bag/ Tin Foil/Envelope (To Protect Sample From Light)
- 7. Venepuncture equipment





# **Explain the procedure**



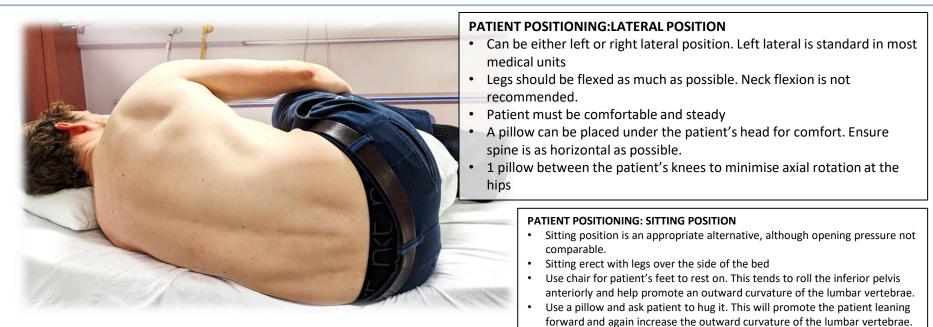
# **Consider contraindications**

# Patient and clinician preparation

- The patient may wish to visit the bathroom prior to positioning
- Ensure clinician to patient position is appropriate and seat available if required. Ideally have the needle at eye level when sitting, as this makes obtaining the correct insertion angle easier
- Remove pager and/or phone
- Ensure your assistant is prepared
- Reassure the patient

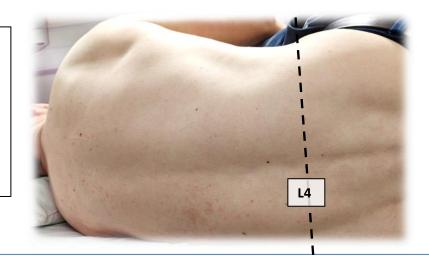


# The procedure: patient positioning and set up



#### **IDENTIFY LANDMARKS**

- Most lumbar punctures are inserted at the L3/4 interspace.
- L2/3 and L4/5 are also suitable in adults.
- Tuffier's line, a line drawn between the posterior iliac crests, corresponds with the body of the L4 vertebra.
- Mark the intended insertion space with an indentation or a marker pen





## The procedure: clinician position and needle control



### **Needle control**

- Insert the introducer using your dominant hand, then hold the introducer with your nondominant hand to maintain control whilst inserting the spinal needle with your dominant hand.
- Rest your non-dominant hand against the patient's back to increase your needle control

### Clinician

- Sit on a chair
- If safe to do so, raise the patient's bed until the spine is at eye level for the clinician. This will make it easier to judge the angle of needle insertion
- Place the procedure trolley nearby, using your assistant to move equipment





# Ensure that the patient is lying on absorbent pads and that clothing has been moved out of the way.

- Wash hands and dry with sterile hand towels.
- Put on sterile gown and gloves.
- Apply antiseptic skin wash.
- Ensure that equipment is not at risk of exposure to antiseptic spray/ liquid drops.
- Allow the chlorhexidine to dry whilst you prepare yourself and your equipment.
- Be vigilant that nothing contaminates the area whilst the chlorhexidine is drying.

## **DRAPE THE PATIENT**

 Remove the adhesive strips from the drape and, without contaminating the field, apply the drape to the patient with the circular cut-out centred on the intended vertebral interspace





Chlorhexidine Gluconate 0.5% w/v in 70% w/v DEB



# The procedure: analgesia & anaesthesia

# LOCAL ANAESTHETIC

- Ask assistant to open and hold vial of lidocaine and draw it up
- Ensure no allergy to lidocaine
- Infiltrate skin with lidocaine. Depending upon the amount of subcutaneous tissue up to 3-4ml of lidocaine 1% or 2% may be appropriate.
- Infiltrate the intended track to the ligamentum flavum, there should be resistance to infiltration once the ligament is reached.

# ANALGESIA

- In patients who are very anxious consider oral analgesia or oral anxiolytics prior to the procedure.
- Note conscious sedation should not be carried out in ward areas without anaesthetic support.



# Maximal lidocaine doses

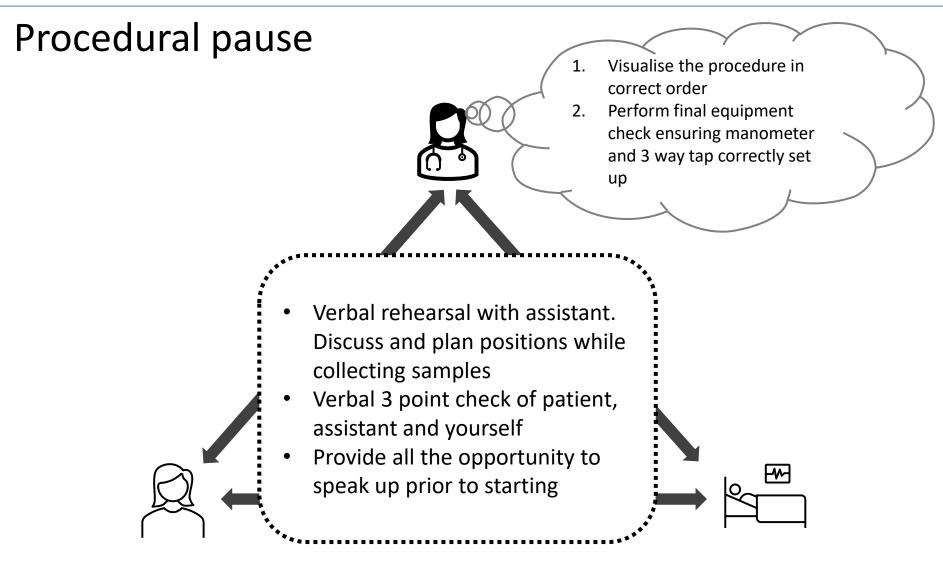
- **4.5mg/kg** in adults and children >12 years previous guidance with different recommendations with/without adrenaline have been removed.
- 3mg/kg in children<12 years
- Not to exceed **200mg** in healthy adults <u>Lidocaine Hydrochloride Injection Summary of Product</u> <u>Characteristics (SmPC) - (emc) (medicines.org.uk)</u>

Example for 60kg patient: Local Anesthetic Dosing Calculator - MDCalc

- Maximal dose = 200mg (290mg by weight but note 200mg max)
- Lidocaine 1% = 1g/100ml
- Maximum dose = 20mL

Formula: patient weight(kg)x 0.45 = maximal lidocaine 1% dose in mL







# Insertion

# **INSERTION OF SPINAL NEEDLE**

Confirm your landmarks and the spinal interspace. Note that an introducer is required for the pencil point needle you will use; this will be included with the spinal needle in its sterile pack.

Watch the associated video for more details about needle technique.

## **OPENING PRESSURE**

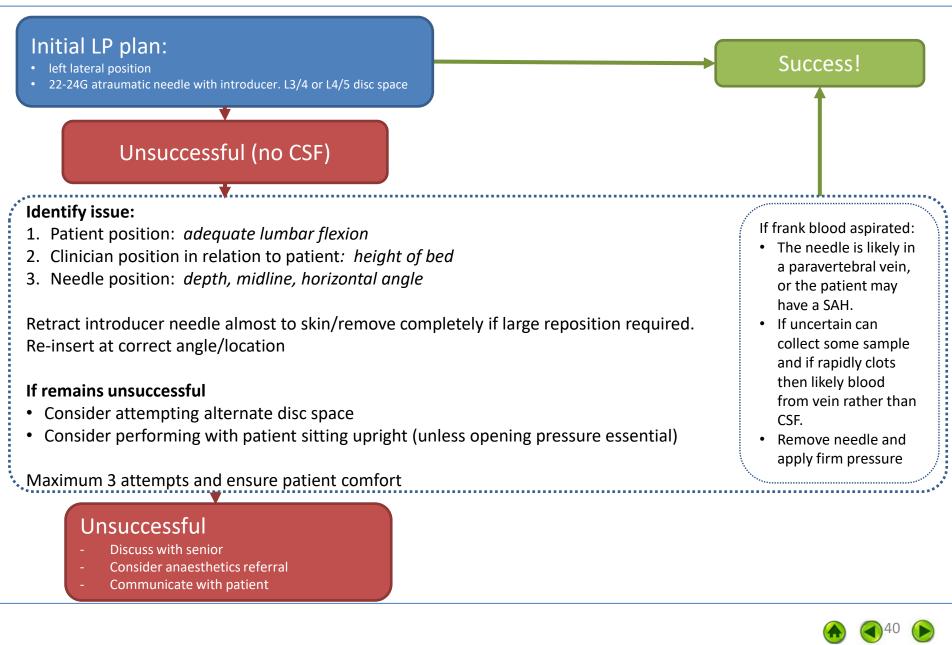
Use the manometer and measure the opening pressure for every patient (95% reference interval for normal pressure is 10-25 cmCSF)







# The procedure: troubleshooting



# **CSF COLLECTION**

- Each drop of CSF is around 0.06mL. 10 drops = 0.6mL
- >2ml (~30 drops) is required for microscopy and culture (MC&S)
  - 10 drops is **not sufficient.**
  - Taking insufficient samples reduces sensitivity, results in some investigations not being performed and may result in the need for a repeat LP
  - Needle size and type does not significantly affect droplet size
- Other investigations may require less volume, details can be found <u>here</u>
- If sample bottles are not externally sterile, the assistant should hold sample collection bottles under LP needle, taking care not to touch anything sterile.
- Ensure the bottle tops are secure.

The safe volume of CSF to take at LP is around 15ml in adults (~250 drops).

Elderly patients may produce reduced volumes







## Biochemistry

- CSF protein, glucose (min volume 0.2ml).
  - Paired serum glucose is essential for interpretation.
  - Ideally send CSF Glucose in yellow top tube (as more stable)
- CSF lactate is a useful test to discriminate bacterial & viral meningitis with reported sensitivity 93% and specificity 96% (if sent prior to antibiotics)<sup>2,22, 23</sup>
- Requests for spectrophotometry to exclude SAH should be made on the last sample of CSF collected (min volume 0.2ml) and need to be protected from light using an envelope or tin foil
- If testing for Oligoclonal bands, paired serum protein essential

# Microbiology/Virology

- Microscopy, culture and sensitivity (MC&S) (volume >2ml, ~30 drops)
- Request CSF viral and bacterial PCR on all possible meningitis/encephalitis CSF samples
- Send EDTA (red top tube) blood sample for bacterial PCR (Meningococcus, Pneumococcus, Haemophilus)
- if TB considered, at least 6ml CSF required<sup>24</sup>
- If CSF antibodies requested, a paired serum sample (brown top tube) is essential (e.g. CNS Lyme)
- Always send blood cultures for patients with suspected CNS infection

## Pathology

- at least 5ml if malignancy considered, usually for cytospin



# Post procedure: Specimen transport and processing

- Ensure correct minimum volume
- The specimens must be labelled correctly and sealed
- Ensure there are sufficient clinical details in TRAK/on request form as this will affect the tests performed
- Ensure timely arrival at the lab and inform Biomedical Scientist (BMS) that microscopy for cell count required. You should know the process for your local site
  - Details on contacting BMS and how to send samples by each site here: <u>Test Directory | Edinburgh and Lothians Laboratory Medicine</u> (edinburghlabmed.co.uk)
- Consider HIV testing in all patients with Meningoencephalitis as per <u>BHIVA/BASHH/BIA Adult HIV Testing guidelines 2020</u>
- Approximately one white cell is permitted for every 500 cells in the CSF cell count

CSF for cell count is an urgent sample<sup>25</sup>

Cell counts decrease rapidly with time, with a drop in reported neutrophil count of 32% at 1 hr and 50% at 2hr.

Leukocyte survival in cerebrospinal fluid. - PMC (nih.gov)



## Post procedure: completion

- 1. Remove needle and apply pressure with gauze until bleeding stops
- 2. Provide advice to the patient about follow up/ symptoms to be aware of such as delayed Post LP Headache
- 3. Label and send CSF samples
- 4. Return bed to usual position
- 5. Apply simple dressing to insertion site and ensure no ongoing bleeding
- 6. Dispose of waste and sharps appropriately
- 7. Restock LP trolley/box if in use.
- 8. Document procedure

# Why documenting LP accurately is important:

- Communicate to colleagues how procedure was done (may affect results e.g. microscopy)
- Medio-legal if complication occurs (e.g. post dural puncture headache, spinal haematoma)
- Documenting which tests were sent helps team review correct results (and act if results not received)
- Documenting appearance of CSF, e.g. if initial blood clears suggests traumatic LP rather than SAH



## Post procedure: documentation

## An example documentation short code from NHS Lothian, entered using "\lp"

LUMBAR PUNCTURE DOCUMENTATION

Patient information leaflet & consent form available here: intranet.lothian.scot.nhs.uk/Directory/neurology/PatientLeaflets/LumbarPunctureleaflet.pdf Information leaflet provided: Y/N/AWI

Indication and procedure explained: Y/N/AWI Risk of side effects explained: Y/N/AWI Consent obtained: Y/N/AWI

Pre-procedure checks completed: Y/N No contra-indications on review: brain imaging; blood tests; allergies; medications: Guidelines on anti-coagulants and anti-platelet agents: intranet.lothian.scot.nhs.uk/Directory/anaestheticsandtheatres/AnaestheticsTheatresDocs/Documents/PreoperativeAssessmentandPerioperativeCare/Warfarin, Direct Oral Anticoagulants and Anti-platelet Medication

Operator/supervisor: Position: lateral/sitting Vertebral level: Aseptic technique: Y/N Local anaesthetic type/dose: Attempts: Needle type/gauge: Opening pressure: Volume CSF removed:

Always send CSF samples for: *Cell count C&S* (*culture & sensitivities*) *Protein Glucose (always send paired serum sample within 4 hours*) *Consider sending CSF samples for: Viral PCR (recommend paired serum HIV test*) *Oligoclonal bands (always send paired serum sample) Spectroscopy/xanthochromia (protect from light and transport by hand, phone the lab if approaching out of hours*) *Cytology (5-10mls) Lactate Mycobacteria TB lx (5-10mls) Mycology fungi lx (5-10mls)* 

Post-procedure blood tests obtained: Y/N Must send: Paired serum glucose (within 4 hours) Consider: HIV test; paired serum oligoclonal bands; bacterial meningitis PCR profile; blood cultures

Post-procedure advice given: Y/N

The clinical test directory can be found here: https://edinburghlabmed.co.uk/TestDirectory/Pages/default To ensure accurate reporting of cell count, do not delay in taking samples to the lab



## Summary

Thank you for completing this pre-learning You are now ready to attend an in-person Mastery Learning Session

By the end of this Mastery pre-learning package you will be able to:

- 1. Describe the indications for performing a lumbar puncture and its use in clinical practice
- 2. Explain the risk assessment, patient safety concerns and contraindications to lumbar puncture
- 3. Describe the potential complications of the procedure and the basic principles of their management.
- 4. Explain the practicalities of performing a lumbar puncture in a safe and structured fashion.
- 5. Recognise your limitations and when to obtain help from a senior clinician.
- 6. Describe the process for ensuring the correct CSF results are requested and obtained.





# **Reference** list



- 1. Grindborg O, Naucler P, Sjolin J *et al.* Adult bacterial meningitis-a quality registry study: earlier treatment and favourable outcome if initial management by infectious diseases physicians. Clin Microbiol Infect Off Publ Eur Soc Clin Microbiol Infect Dis 2015;21(6):560e6.
- 2. McGill, F. *et al.* (2016) 'The UK joint specialist societies guideline on the diagnosis and management of acute meningitis and meningococcal sepsis in immunocompetent adults', *Journal of Infection*, 72(4), pp. 405–438. doi: 10.1016/j.jinf.2016.01.007.
- 3. Proulx N, Frechette D, Toye B et al. Delays in the administration of antibiotics are associated with mortality from acute bacterial meningitis. QJM 2005;98:291e8.
- 4. Auburtin M, Wolff M, Charpentier J, *et al.* Detrimental role of delayed antibiotic administration and penicillin-nonsusceptible strains in adult intensive care unit patients with pneumococcal meningitis: the PNEUMOREA prospective multicenter study. Crit Care Med 2006;34:2758e65.
- 5. Koster-Rasmussen R, Korshin A, Meyer CN. Antibiotic treatment delay and outcome in acute bacterial meningitis. J Infect 2008;57(6):449e54.
- 6. Dodd, K. C. *et al.* (2018) 'Periprocedural antithrombotic management for lumbar puncture: Association of British Neurologists clinical guideline', *Practical Neurology*, 18(6), pp. 436–446. doi: 10.1136/practneurol-2017-001820.
- 7. Nath S, et al. Atraumatic versus conventional lumbar puncture needles: a systematic review and meta-analysis. Lancet. 2018 Mar 24;391(10126):1197-1204.
- 8. NAP3: Major Complications of Central Neuraxial Block in United Kingdom, Available from: https://www.nationalauditprojects.org.uk/NAP3\_home?newsid=464 [Accessed on 28th June 2022]
- 9. Joffe AR. Lumbar puncture and brain herniation in acute bacterial meningitis: a review. J Intensive Care Med. 2007;22:194-207. DOI: 10.1177/0885066607299516
- 10. Tourtellotte WW, Henderson WG, Tucker RP, et al. A randomized, double-blind clinical trial comparing the 22 versus 26 gauge needle in the production of the post-lumbar puncture syndrome in normal individuals. Headache J Head Face Pain 1972;12(2):73e8.
- 11. Carson D, Serpell M. Choosing the best needle for diagnostic lumbar puncture. Neurol 1996;47(1):33e7.
- 12. Thomas SR, Jamieson DR, Muir K. Randomised controlled trial of atraumatic versus standard needles for diagnostic lumbar puncture. BMJ 2000;321:986e90.
- 13. Norris MC, Leighton BL, DeSimone CA. Needle bevel direction and headache after inadvertent dural puncture. Anesthesiology 1989;70:729e31.
- 14. Richman JM, Joe EM, Cohen SR, et al. Bevel direction and postdural puncture headache. Neurologist 2006;12:224e8.
- 15. Strupp M, Brandt T, Muller A. Incidence of postlumbar puncture syndrome reduced by reinserting the stylet: a randomized prospective study of 600 patients. J Neurol 1998;245:589e92.
- 16. MacArthur C, Lewis M, Know EG. Accidental dural puncture in obstetric patients and long term symptoms. BMJ 1993;306: 883e5.
- 17. Kuntz KM, Kokmen E, Stevens JC, Miller P, Offord KP, Ho MM. Post-lumbar puncture headaches: experience in 501 consecutive procedures. Neurology 1992;42(10):1884e7.
- 18. Sung RK, Hyun SC, Mi JY, Jung HH, Kwang JC, Sun JC. No effect of recumbency duration on the occurrence of postlumbar puncture headache with a 22G cutting needle. BMC Neurol 2012;12(1):1e5.
- 19. Arevalo-Rodriguez, Ingrid, et al. "Posture and fluids for preventing post-dural puncture headache." Cochrane Database of Systematic Reviews 3 (2016).
- 20. Rochwerg B, Almenawer S A, Siemieniuk R A C, Vandvik P O, Agoritsas T, Lytvyn L et al. Atraumatic (pencil-point) versus conventional needles for lumbar puncture: a clinical practice guideline BMJ 2018; 361 :k1920 doi:10.1136/bmj.k1920
- 21. Checketts, M.R. (2012), Wash & go but with what? Skin antiseptic solutions for central neuraxial block. Anaesthesia, 67: 819-822. https://doi.org/10.1111/j.1365-2044.2012.07263.x
- 22. Huy, N.T., Thao, N.T., Diep, D.T. et al. Cerebrospinal fluid lactate concentration to distinguish bacterial from aseptic meningitis: a systemic review and meta-analysis. Crit Care 14, R240 (2010). https://doi.org/10.1186/cc9395
- 23. Kristian Buch, Jacob Bodilsen, Andreas Knudsen, et al. for the Danish Study Group for Infections in the Brain (2018) Cerebrospinal fluid lactate as a marker to differentiate between communityacquired acute bacterial meningitis and aseptic meningitis/encephalitis in adults: a Danish prospective observational cohort study, Infectious Diseases, 50:7, 514-521, DOI: 10.1080/23744235.2018.1441539
- 24. Thwaites GE, Chau TT, Farrar JJ. Improving the bacteriological diagnosis of tuberculous meningitis. J Clin Microbiol. 2004 Jan;42(1):378-9. doi: 10.1128/JCM.42.1.378-379.2004. PMID: 14715783; PMCID: PMC321694.
- 25. Steele RW, Marmer DJ, O'Brien MD, et al. Leukocyte survival in cerebrospinal fluid. J Clin Microbiol. 1986 May;23(5):965-6. doi: 10.1128/jcm.23.5.965-966.1986. PMID: 3711287; PMCID: PMC268763.
- 26. van Dongen RM, Onderwater GLJ, Pelzer N, *et al.* The effect of needle size on cerebrospinal fluid collection time and post-dural puncture headache: A retrospective cohort study. Headache. 2021 Feb;61(2):329-334. doi: 10.1111/head.14046. Epub 2021 Jan 16. PMID: 33452678; PMCID: PMC7985863.
- 27. Thwaites G, Fisher M, Hemingway C, *et al.* British Infection Society. British Infection Society guidelines for the diagnosis and treatment of tuberculosis of the central nervous system in adults and children. J Infect. 2009 Sep;59(3):167-87. doi: 10.1016/j.jinf.2009.06.011. Epub 2009 Jul 4. PMID: 19643501.

