

PAJUNK®

IntraLong

Regional Anesthesia



Instructions for use

 These instructions for use were translated into the following languages: DE, EN, FR, IT, ES, PT, NL, DA, SV, EL, BG, ET, HR, LV, LT, PL, RO, SK, SL, CS, HU. The translations can be downloaded from our website: eifu.pajunk.com.

Special notice

 Please read the following information and operating instructions carefully!
 **Caution:** Federal law restricts this device to sale by or on the order of a physician. The device may only be used by qualified medical staff in accordance with these user instructions.

PAJUNK® does not recommend any particular treatment method. Professional medical staff are responsible for the way in which the device is used and for patient selection.

In addition to these instruction for use, the relevant information also applies according to the corresponding specialist literature and current state of the art and knowledge.

Failure to comply with the instructions for use invalidates the warranty and puts patient safety at risk.

If used in combination with other devices, it is essential that the compatibility information and user instructions for these other devices are taken into account. A decision regarding the combined use of devices from different manufacturers (where they do not constitute treatment units) is the responsibility of the user.

 The device must not be used under any circumstances if there are good reasons to suspect incompleteness, damage or loss of sterility.

 Only devices in perfect condition, which are within the sterile expiry date marked on the label, in undamaged packaging, may be used.

The “Summary of Safety and Performance acc. to EU-2017-745 (SSCP)” is available from EUDAMED.

Device description / compatibility

 Please see the current declaration of conformity for product numbers and the scope of these instructions for use.

IntraLong is provided by PAJUNK® in convenient kits consisting of:

- Cannula: SPROTTE® SPECIAL tip, retaining plate
- Catheter (25-27G with stylet, fully radiopaque) in catheter container
- Clamping Adapter
- Locking cap
- Bacterial filter: 0.2 µm
- Syringe

Hub connectivity: LUER

The exact composition may be gathered from the label.

Intended use

Puncture, access to the target area, aspiration, injection, catheter placement

The catheters are intended to remain in the target area (subarachnoid space) and constantly deliver a local anaesthetic from an external source.

 *Indwelling time for the continuous system: 7 days (168h)*

 *PAJUNK® cannulas can also be introduced into the body under ultrasound, fluoroscopic or CT guidance.*

Target user group

Medical specialist staff only

Target patient population

Adults and children. Professional medical staff are responsible for patient selection.

Indications

Continuous spinal block for surgical anaesthesia, obstetrical analgesia, post-operative analgesia and treatment of chronic pain as well as a complement to general anaesthesia.

Contraindications

Device-specific contraindications

 *Under no circumstances is the device to be used in the event of known material incompatibilities and/or known interactions.*

Clinical contraindications

Absolute contraindications:

- Patient refusal
- Poorly controlled bleeding diathesis or anticoagulation (coagulation disorders)
- Systemic infection (sepsis/bacteraemia)
- Local infection at the injection site
- Local malignancy at the injection site
- Weakened immune system
- Strong, de-compensated hypovolaemia, shock
- Uncontrolled diabetes mellitus

Relative contraindications:

- Specific neurological disorders
- Specific cardiovascular disorders
- Allergic reaction/hypersensitivity to the administered agents (contrast, anaesthetic or corticosteroid)
- Severe deformations of the spine, arthritis, osteoporosis, spinal disc herniation or condition after spinal disc surgery
- Condition after spinal fusion, spinal metastasis

- Recent consumption of non-steroidal anti-inflammatory medications
- Unexperienced user

Complications

Device-specific complications:

Cannula: Cannula bending, breakage, occlusion, leakage at the cannula hub

Catheter: Catheter breakage, catheter shearing, catheter bending, catheter knotting, reduced/absence of flow (occlusion), catheter disconnection.

Procedure-specific complications

Cannula: Undesirable positioning of the cannula (e.g. intravascular, intraneural, etc.), repeated puncture/redirection of the cannula, failed procedure.

Catheter:

During placement:

Inability to place catheter tip within subarachnoid space (results in catheter knotting or shearing on the introduction cannula tip), accidental intravascular catheter placement, difficulty in advancing the catheter (may result in catheter kinking).

During application:

- Technical problems resulting in the premature discontinuation of analgesia due to catheter disconnection, catheter obstructions (occlusion); leakage at the catheter exit site.
- Premature discontinuation of analgesia due to catheter-related infections
- Catheter migration

During removal:

Resistance when removing the catheter resulting in catheter breakage.

Clinical complications

- Local and systemic infections
- Neuronal damage (during cannula placement, which may result in temporary increase in pain, temporary motor weakness, transient back or extremity pain, numbness and/or tingling, paraplegia)
- Accidental vascular punctures with corresponding complications (vascular lesions, bleeding/haematoma, vasovagal reactions, intravascular injection, etc.)
- Intra-arterial injection (direct injection into the spinal cord, vertebral artery or radicular artery; including spinal cord infarct, epidural haematoma and brainstem haemorrhage, neurological events, vascular complications, thrombosis or thromboembolism)
- Puncture of the dura with corresponding complications
 - *Dura puncture and liquor loss:* post-spinal headache or backache, nausea, vomiting, neurological damage, epidural haematoma, epidural abscess
 - *Anaesthetic in the subarachnoid space:* circulatory disorders, decrease of the body temperature, urinary retention, respiratory side effects and complications, extremities weakness, total spinal anaesthesia, cauda

equina syndrome

- Toxicity of local anaesthetic

 *Users must inform patients of complications typically associated with the procedure.*

 *If complications occur while using the device, follow the protocols of your organisation. If this does not resolve the complications, or if they are regarded as serious or untreatable, carefully stop the procedure and remove invasive device components from the patient.*

Warnings

 *for sterile device:*

This is a disposable medical device for use on one patient only!

 *This device must not be re-used under any circumstances!*

 *This device must not be re-sterilised under any circumstances!*

The materials used in the manufacture of this device are not suitable for reprocessing or re-sterilisation.

This device is not designed to be reprocessed or re-sterilised.

 **Unauthorised re-use or reprocessing**

- can cause the device to lose the essential performance properties intended by the manufacturer.
- leads to a significant risk of cross-infection/contamination as a result of potentially inadequate processing methods.
- may cause the device to lose functional properties.
- may cause materials to break down and lead to endotoxic reactions caused by the residues.

 *for puncture:*

1. Take care to use devices of suitable dimensions (diameter, length), especially when treating obese patients and children.
2. Only perform the puncture (even when removing the cannula) with the stylet in place.
3. To avoid bending or breaking of the cannula, never apply excessive force to the cannula.
4. In case of unexpected bone contact, slightly withdraw the cannula and change its direction.
5. Repeated bone contact will damage the cannula tip. Under no circumstances should you continue to use a cannula damaged in this manner. In case of previous bone contact, remove the cannula in one step.

 *for catheter placement and removal:*

1. Immediately before use, check that the catheter will pass through the cannula.
2. The tip of the cannula can be damaged by bone contact during insertion. If a catheter is passed through a cannula that is damaged in this way, it can itself become damaged. If this happens, use a new cannula.
3. Once the catheter has left the tip of the cannula, do not retract the catheter as there is a risk of shearing.
4. If blood is visible in the catheter return window or in the piston chamber of the syringe, remove the catheter and reattempt puncture. The catheter was incorrectly positioned.
5. If the procedure is interrupted, remove the catheter and the cannula together if possible.
6. If flow is impeded, check the locking mechanism of the Clamping Adapter.
7. When using catheters with a closed tip and lateral openings, extend the catheter at least 15 mm (no more than 50 mm) beyond the tip of the cannula to ensure unimpeded injection.
8. Never insert the catheter more than 50 mm. It is more likely to become knotted if it is inserted more than 50 mm.
9. Ensure that the catheter is not kinked on fixing.
10. Be sure to check the connection between the catheter and the infusion devices regularly.
11. Do not tug the catheter or pull it sharply when removing it from the patient.
12. Do not exert excessive force when removing the catheter. Do not continue to pull the catheter if it starts to stretch too much.
13. If you detect resistance while removing the catheter, do not withdraw it any further. If necessary, reposition the patient so as to enlarge the gap between the vertebrae. Then try to withdraw the catheter again. If this is still difficult, investigate with fluoroscopy or an X-ray before taking any further action.
14. After removing the catheter, check the distal tip to see whether it is complete. The tip should be intact. Only in this case can you be sure that the entire catheter has been removed.

 *for injection:*

1. Always ensure that the injection site is aseptic.
2. Do not administer drugs that are not indicated for the intended use.
3. Aspirate before the injection of medication. If you observe blood in the cylinder of the syringe, then the cannula has been introduced improperly. **STOP THE PROCEDURE.**
4. Be sure to regularly check the connection between the cannula/catheter and the infusion device.

 for use with other compatible products:

1. When using multiple components, familiarise yourself with their operation before use by checking connections and passages (cannulas, adapters).
2. When connecting the catheter to the Clamping Adapter, always make sure that the catheter is fully inserted into the Clamping Adapter as far as the stop (at least as far as the orientation mark). Never preflush before making the connection.
3. Disinfectants based on or containing alcohol can damage the filter.
4. The locking cap must be screwed on before you disinfect the filter.

 further warning indications:

1.  Caution: Sharp object warning. The device or device components may, depending on the type of tip, have sharp edges or tips. Various infectious pathogens can be transmitted if a stab wound occurs. The most relevant ones in practice are the human immunodeficiency virus (HIV), the hepatitis B virus (HBV) and the hepatitis C virus (HCV).
2. You must routinely take general precautions for handling blood and body fluids when using and disposing of the device, due to the risk of contact with blood-borne pathogens.
3. Please note that the continued use of a device of the same type must be assessed cumulatively as described in the legislation on medical devices, even after the device has been exchanged or replaced.
4. Avoid build-up of fluid film between the catheter and Clamping Adapter (e.g. through fluids on gloves). Fluids on the proximal end of the catheter can affect the holding force and result in disconnections and/or leakage.

Sequence of use

Placement of the spinal cannula

1. Perform skin disinfection and cover area to be punctured with a sterile fenestrated surgical drape (aperture drape).
2. Administer a local anaesthetic.
3. If necessary, perform a perforating incision of the area to be punctured (lancet, or similar).
4. Advance the cannula up to the subarachnoid space.
5. Retract the stylet from the cannula.
6. Identify the subarachnoid space through corresponding CSF reflux.

Placement of the spinal catheter

1. Place the catheter container on the cannula hub.
2. Push the catheter with the marked end into the target area until it has reached the required depth. Take the catheter out of the catheter container and withdraw the container until the stylet has also been completely pulled out of the catheter.
3. The correct catheter position must be checked by means of CSF reflux.
4. Once it is in place, remove the cannula via the catheter. Hold the catheter tightly with the other hand if necessary.
5. After removing the cannula, connect the catheter to the Clamping Adapter.
6. Fill the filter with the anaesthetic solution to be used at the beginning of the anaesthesia/analgesia to compensate for the dead volume (the filling volume of the filter is approximately 0.35 ml).
7. Connect the Clamping Adapter to the hub of the filter.
8. Fill the syringe with the selected anaesthetic or analgesic and connect it to the filter hub. The catheter system is now ready for the injection (depending on age and weight of the patient, as well as on the type of the intervention and the composition of the anaesthetic).
9. Secure the catheter at the exit site using the optionally supplied FixoCath.

Use and storage conditions

	Temperature limit	+10 °C to +30 °C
	Humidity limitation	20 % to 65 %
	Keep away from sunlight	
	Keep dry	

General information

The devices are manufactured in accordance with globally applicable guidelines for hazardous substances.

 Non-pyrogenic

 *Any serious incident that occurred while using the device should be reported to the manufacturer and the corresponding authorities of the country the user and/or patient are residing in.*

 PAJUNK® GmbH Medizintechnologie, Karl-Hall-Strasse 1, 78187 Geisingen, Germany.

Key to symbols used in labelling



Manufacturer



Use-by date



Item number



Sterilized using ethylene oxide



Do not re-sterilise



Do not use if package is damaged



Keep dry



Humidity limitation



Do not re-use



Caution



Date of manufacture



Batch code



Keep away from sunlight



Temperature limit



Consult instructions for use



Single sterile barrier system



Non-pyrogenic



Dispensing with prescription only (the device may only be used by qualified medical staff for the intended purpose.)



Advice



Information



“CE conformity marking” or “CE marking” = this marking shows that a device is in conformity with the applicable requirements as set out in the Medical Device Regulation or other European Union legislation on its affixing.



Sharp object warning



Does not contain phthalates



Does not contain natural rubber latex



Quantity



Translation



Medical device



Unique device identification



Single sterile barrier system with protective packaging outside



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