Instructions for use

CE 2862

[Product Name]

Anesthesia Needle

[Intended Use]

AN-S is intended for administration of local anesthetics into subarachnoid cavity. AN-E is intended for administration of local anesthetics into epidural space.

[Type and Specification]

Type	Specification
AN- S	20G,22G,25G,26G,27G
AN-E	16G,17G,18G,19G

[Indication]

Lower abdominal surgery, anal and perineal surgery, pelvic surgery, lower limb surgery and analgesia.

[Contraindications]

Includes diseases or conditions such as:
1. Patient refusal: psychosis disorders, severe neuroses, and uncooperative child;
2. Severe hypovolemia;
3. Disorders of Coagulation;
4. Infection at the puncture site;
5. Diseases of the central nervous system;
6. Spinal trauma or a history of severe back pain or unknown spinal nerve compression disorders;
7. Systemic infection.

[Patient target group]

Adult and child.

[Intended users]

Professional medical staff

[Use Method]

a) Read the instructions for use carefully before use.

b) Place patients in proper position.

c) Check whether the package is damaged, and it is forbidden to use if it is damaged; Open the single package, wear gloves according to the aseptic operation specification, and check whether the contained device is complete.

d) Disinfect the puncture site and perform subcutaneous local anesthesia according to routine procedures.

e) Epidural needle is used according to the method of epidural space puncture.

f) Spinal needle is used according to the method of subarachnoid cavity puncture.

[Warnings and Precautions]

1. The use is prohibited if one of the following situations occurs: (1) the single package is damaged; (2) the product is expired; (3) Deformation or damage of accessories.
2. This device should be used by trained professional medical staff.
3. Be gentle during puncture process, do not operate roughly.
4. If excessive resistance is met during needle insertion, do not force the needle as damage may occur.
5. If the spinal or epidural needle is bent during use, it is forbidden to straighten the needle and continue to use it.
6. This product is for single use. Please destroy it after use, the destruction procedures shall be handled harmlessly by qualified or authorized institutions according to the local relevant regulations.
7. The product has been sterilized by Ethylene oxide with validity for 5 years. Re-sterilize and/or reuse the product can potentially result in compromised device performance and increased risk of inappropriate destabilization and cross contamination.
8. Please inform the manufacturer in case of any incidents related to the device occur and report to the competent authority of the Member State in which the incidents occurred, if applicable.

[Side effects]

Cardiovascular complications, respiratory complications, Total spinal anesthesia, spinal nerve anomalous block, nausea and vomiting, urinary retention, intradural hematoma, hemorrhage, infection, dural puncture headache, neurological injury, spinal cord ischemic injury and anterior spinal artery syndrome.

[Shelf-life]

5 years

[Sterilization Method]

Ethylene oxide

[Clinical benefits]

Establish a channel to the epidural space or subarachnoid cavity for the patient, so that the anesthesia drugs can

reach the epidural space or subarachnoid cavity.

[Performance characteristics]

The conical connector of the needle hub meet the requirements of ISO 80369-6:2016.

[Flow rate]

Gauge	Flow rate (ml/min)
25G	≥6
27G	≥3

[Storage and Transport Conditions]

1. No heavy pressure, direct sunlight, rain or snow dipping, so as not to damage the device.
2. Handle with care when transporting to avoid violent collision.
3. Keep away from fire and heat sources in case that the device might be out of shape.
4. This product should be stored in a non-corrosive gas, well-ventilated and clean environment.

[Production Date]

See on the package.

[Symbol Explanation]

	Manufacturer	CN	Date of manufacture, Country of manufacture
EC REP	Authorized representative in the European Community/ European Union	LOT	Batch code
	Use-by date		Do not use if package is damaged and consult instructions for use
	Consult instructions for use or consult electronic instructions for use		Caution
	Do not resterilize, Indicates a medical device that is not to be resterilized		Do not re-use
	Single sterile barrier system	STERILE EO	Sterilized using ethylene oxide
MD	Medical device	UDI	Unique device identifier
CE 2862	CE Marking	REF	Catalogue number
	Keep away from sunlight		Keep dry

[Additional Information]

The SSCP of this product will be available in the European database on medical devices (Eudamed), where it is linked to the Basic UDI-DI (6938119333359JF, 693811933360HY). The URL to the Eudamed public website is: https://ec.europa.eu/tools/eudamed
For a patient/user/third party in the European Union and in countries with identical regulatory regime (Regulation 2017/745/EU on Medical Devices); if during the use of this device or as a result of its use, a serious incident has occurred, please report it to the manufacturer and/or its authorized representative and to your national authority. The contacts of national competent authorities (Vigilance Contact Points) and further information can be found on the following European Commission website: https://ec.europa.eu/growth/sectors/medical-devices/contacts_en

[Manufacturer]
Henan Tuoren Medical Device Co., Ltd.
Address: Weiyuan Industrial Zone, Menggang, Changyuan, 453400 Henan Province, P.R. China
Tel.: +86 0373-8605444
Fax: +86 0373-8605321
E-mail: info@etuoren.com
Website for eIFU: en.tuoren.com/ifu

[EU Representative]
MedNet EC-REP CIII GmbH
Address: Borkstrasse 10, 48163 Münster, Germany

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