

DTR.M.049 FLUSHING NEEDLES

Instructions for Use

REF

HFN2001, HFN2081, TAC2003, TAC20LC, TAC20LO,
TAC20MO, TAC20SC, TAC20SO, TAC20XLC, TAC20XXLC





DESCRIPTION

The Flushing Needles are disposable devices used to probe and flush heparin and/or saline into a vein or artery during surgical procedures.

MATERIALS OF CONSTRUCTION

Product Component	Details
Flushing Needles	Stainless Steel

INTENDED USE

The Flushing Needles are intended to provide an atraumatic way to access small blood vessels for probing and flushing solutions. For this purpose, the instrument features a small needle that is ideal to probe and flush heparin and/or saline into arteries and veins. It is fitted with a profiled nozzle to avoid reflux during manual flushing.

INTENDED USER

The device is intended to be used by appropriately trained surgeons.

INDICATIONS, CONTRAINDICATIONS AND PATIENT SELECTION FACTORS

The devices are indicated for probing and flushing with heparin and/or saline into a vein or artery.

Do not use with in direct contact with the heart, central circulatory (arteriae pulmonales, aorta ascendens, arcus aortae, aorta descendens to the bifurcatio aortae, arteriae coronariae, arteria carotis communis, arteria carotis externa, arteria carotis interna, arteriae cerebrales, truncus brachiocephalicus, venae cordis, venae pulmonales, vena cava superior and vena cava inferior) and central nervous system (brain, meninges and spinal cord).

These devices are made of Stainless Steel, which may cause an allergic reaction in patients with sensitivity to Nickel. Surgeons should risk assess the use of the device in relation to the medical benefit of the procedure and take necessary precautions with patients with known sensitivity to Nickel.

PATIENT POPULATION

Anyone of any age in the human population as determined by a suitably qualified surgeon.

WARNINGS AND PRECAUTIONS

- This device is supplied **STERILE** and ready to use.
- The device is for **SINGLE USE ONLY**. Do **NOT** re-sterilise or re-use.
- Do not use if the packaging has been opened or damaged.
- If the device accidentally becomes contaminated before use, do not use and dispose of accordingly.

SHELF LIFE AND STERILITY

- Instruments are supplied in a sterile condition, packaged individually. The instruments are sterilized by exposure to Ethylene Oxide indicated by the symbol on the label. They remain sterile as long as the package integrity has not been violated.
- Packaging must be inspected before use. Do not use any component from an opened or damaged package. The expiration date is printed on the label. Do not use instruments after the expiration date. The shelf life is 5 years.

DEVICE LIFETIME

Devices are single use transient devices. Therefore, the device lifetime in use is less than 60 minutes continuous use.

STORAGE AND HANDLING

- Sterile, packaged instruments should be stored in a designated, limited access area that is well ventilated and provides protection from dust, moisture, insects, vermin, sunlight and temperature/humidity extremes.
- Care must be exercised in handling of wrapped cases or individual instruments to prevent damage to the sterile barrier.
- Sterile instrument packages should be carefully examined prior to opening to ensure that package integrity has not been compromised.

TRANSPORTATION

The instruments are supplied in their designated packaging. This packaging ensures that every instrument is kept in a way that they do not receive any damage and that their functionality is preserved during transportation.

OPERATIONAL USE

Small opening is made into the vein just below the clamp and flushing needle is inserted and attached to a syringe via luer lock ready for flushing.

DEVICE DISPOSAL

After use the device should be disposed of in accordance with the healthcare settings disposal procedure. The product comes into contact with bodily fluids, which can be contaminated. Care should be taken in the handling and disposal of the device after use to prevent contamination.






The Flushing Needles do not contain:















- Substances of human origin
- Animal tissues
- Dangerous or hazardous substances
- Radioactive material
- Any substance that may be a Medicinal Product as defined in Article I of Directive 2001/83/EC

REPORTING PROBLEMS

Healthcare professionals, users and patients should report any suspected serious incident related to the device by informing the manufacturer at www.dtrmedical.com/contactandthe competent authority, ministry of health, or delegated agency in which the suspected serious incident has occurred.

SYMBOL REFERENCE TABLE

	Indicates the manufacturer's catalog number so that the medical device can be identified.
	Indicates the manufacturer's batch code so that the batch or lot can be identified.
	Indicates the authorized representative in the European Community/ European Union.
	Indicates the packaged quantity.
	Authorized Representative in Switzerland.

	Indicates the date after which the medical device is not to be used.
	Indicates a medical device that has been sterilized using ethylene oxide.
	Indicates a medical device that is not to be re-sterilized.
	Indicates a medical device that is intended for one single use only.
	Keep away from sunlight.
	Keep dry.
	Indicates the item is a medical device.
	Indicates Medical Device Manufacturer.
	Indicates a medical device that should not be used if the package has been damaged or opened and that the user should consult the instructions for use for additional information.
	Indicates the need for the user to consult the instructions for use for important, cautionary information such as warnings and precautions that cannot, for a variety of reasons, be presented on the medical device itself.
	Indicates the need for the user to consult the instructions for use.
GB	To identify the country of manufacture of products (Made in the United Kingdom).
	Indicates the entity distributing the medical device into the locale.
	Indicates a carrier that contains Unique Device Identifier Information.
2797	Indicates that the product conforms to EU Regulation 2017/745 and meets applicable health, safety and environmental requirements. If the mark is accompanied by a number, conformity is verified.
	Indicates that a product conforms to the Medical Device Regulations 2002 (SI 618) as subsequently amended by the EU Exit Regulations of 2019 (SI 791) and 202 (SI 1478) and meets applicable health, safety, and environmental requirements. If the mark is accompanied by a number, conformity is verified.



DTR MEDICAL® BY



DTR Medical Ltd, 17 Clarion Court,
Clarion Close, Swansea, SA6 8RF, UK
T: +44(0) 1792 797910
www.dtrmedical.com