

DTR.M.042 TISSUE FORCEPS

Instructions for Use

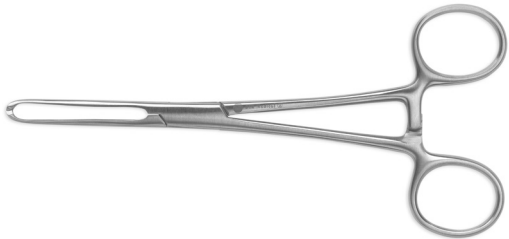
REF

ATF2341, KMF1025

KMF1025
Keele
Mucosectomy
Forceps



ATF2341
Allis Tissue
Forceps





DESCRIPTION

The Tissue Forceps are hinged devices designed for grasping and holding tissue.

The Allis Tissue Forceps are scissor-like forceps, whereas the Keele Mucosectomy Forceps are tweezer-like.

MATERIALS OF CONSTRUCTION

Product Component	Details
Tissue Forceps	410 Stainless Steel

INTENDED USE

The Allis Tissue Forceps are intended to grasp, hold and manipulate soft tissue, organs, skin and underlying tissue.

The Keele Mucosectomy Forceps are intended for control, stabilisation and manipulation of soft tissue, the tongue and carcinoma removal during transoral mucosectomy procedures.

INTENDED USER

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

INDICATIONS, CONTRAINDICATIONS AND PATIENT SELECTION FACTORS

The Allis Tissue Forceps are indicated for general surgical procedures to grasp, hold and manipulate soft tissue, organs, skin and underlying tissue.

The Keele Mucosectomy Forceps are indicated for use during transoral mucosectomy procedures for control, stabilisation and manipulation of soft tissue, the tongue and carcinoma removal.

These devices are made of Stainless Steel, which may cause an allergic reaction in patients with sensitivity to Nickel. The User 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.
- Keele Mucosectomy Forceps are sharp. Caution should be exercised at all times when handling and using these devices.

SHELF LIFE AND STERILITY

The devices are sterilised by exposure to Ethylene Oxide, as indicated by the symbol on the label. They remain sterile as long as the package integrity has not been compromised.

The expiration date is printed on the label. Do not use devices after the expiration date. The shelf life is 5 years.

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

STORAGE AND HANDLING

Sterile, packaged devices 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 individual devices to prevent damage to the sterile barrier.

Sterile device packages should be carefully examined prior to opening to ensure that package integrity has not been compromised.

TRANSPORTATION

The devices are supplied in their designated packaging. This packaging

ensures that every device is kept in a way that they do not receive any damage and that their functionality is preserved during transportation.

OPERATIONAL USE

This is a standard device, for operational use please refer to indications and intended use.

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.

Keele Mucosectomy Forceps are sharp and must be placed in a secure puncture-resistant bin (conforming to appropriate regulations) which is suitable for incineration.
















The Tissue Forceps 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/contact and the competent authority, ministry of health, or delegated agency in which the suspected serious incident has occurred.

SYMBOL REFERENCE TABLE

	Indicates the manufacturer's catalogue 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 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 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 a medical device that is intended for one single use only.
	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.
	Indicates Medical Device Manufacturer.
	To identify the country of manufacture of products (Made in the United Kingdom).
	Indicates the packaged quantity.
	Indicates the entity distributing the medical device into the locale.
	Indicates the item is a medical device.
	Indicates a carrier that contains Unique Device Identifier Information.

	<p>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.</p>
	<p>Indicates a medical device that needs protection from light sources.</p>
	<p>Indicates a medical device that needs to be protected from moisture.</p>
	<p>Caution: US Federal law restricts this device to sale by or on the order of a physician.</p>



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