

1 Instructions for Use

Products:

E 01 2107 PointerTube Keat
E 01 2108 PointerTube Kennedy

are accessories of the Cube Navigation System

Trade Name: Cube Navigation System
Common Name of Device: Image guided surgery system

Manufacturer:

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For US market only. This document is intended to provide information to an audience of the US.

Explanation of symbols



Manufacturer / Manufacturing date



Reference number/
Order number



Serial number



To ensure safety, follow the instructions for use



MR UNSAFE. The system must not be used in a Magnetic Resonance Environment. It contains ferromagnetic parts and pose a clear threat to persons and equipment in the magnetic room.



Note to the user to adhere to and consult the referenced Instructions for Use of other system components or 'Application Notes'



Attention
Follow the supporting documentation



Indicates the temperature limits to which the medical device can be safely exposed



Indicates the humidity limits to which the medical device can be safely exposed



Application part type BF



This is the general warning sign. It is used to alert the user to potential hazards. All safety messages that follow this sign shall be obeyed to avoid possible harm.



U.S. Federal law restricts this device to sale by or on the order of a physician

1. General information

The products PointerTube Keat and PointerTube Kennedy are variants of a navigated suction instrument for image guided surgical procedures and are referred to as *'instruments'* below.

The instruments are part of the Cube Navigation System which is designed to be used with the Cube Navigation System. The instruments are accessories to the Cube Navigation System produced by Fiagon GmbH.



CAUTION: Read the instructions for use carefully before using the instrument.

CAUTION: Before using it, the instrument must be reprocessed according to the reprocessing instructions. All instruments are delivered in NON-STERILE condition.

The instruments can be reprocessed and used 10 times. The Cube Navigation System indicates the remaining number of uses.

2. Indications for Use / Field of application



The application of the instrument is limited to the indications for use described here.

The instruments are intended as an aid for precisely locating anatomical structures in either open or percutaneous procedures. They are indicated for use with the Cube Navigation System using electromagnetic navigation.

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It is indicated for any medical condition in which the use of stereotactic surgery may be appropriate, and where reference to a rigid anatomical structure in the field of ENT surgery, such as the paranasal sinuses, mastoid anatomy, can be identified relative to a CT or MR based model of the anatomy.

Example procedures include, but are not limited to:

ENT Procedures;

- Transsphenoidal access procedures.
- Intranasal procedures.
- Sinus procedures, such as Maxillary antrostomies, Ethmoidectomies, Sphenoidotomies/Sphenoid explorations, Turbinate resections, and Frontal sinusotomies.
- Skull base procedures for ENT access

The navigation system and its accessories are intended for use by healthcare professionals only. In addition, the users receive a training. The operator, i.e., the person or facility that is responsible for the use and service of the system, must ensure that all users of the system receive an adequate introduction into the system in accordance with valid laws and regulations. An operator is everyone who uses the system.

The instrument as a part of the navigation system serves as a tactile instrument. The position of the tip of the instrument and possibly the orientation of the axis are displayed in the image data. The instructions for use of the navigation system describe the proper use of the entire system in detail.

3 Instructions for Use



The instrument may only be used if the safety instructions of the navigation system and other connected devices have been followed.

The navigation only works in connection with the navigation system of Fiagon GmbH.



Contraindications

CAUTION: The instrument should not be used if the following contraindications exist:

The instrument should not be re-processed and reused after it has been used with patients with suspected Creutzfeldt-Jakob disease. There is a risk of disease transmission, dispose the instrument after the operation.

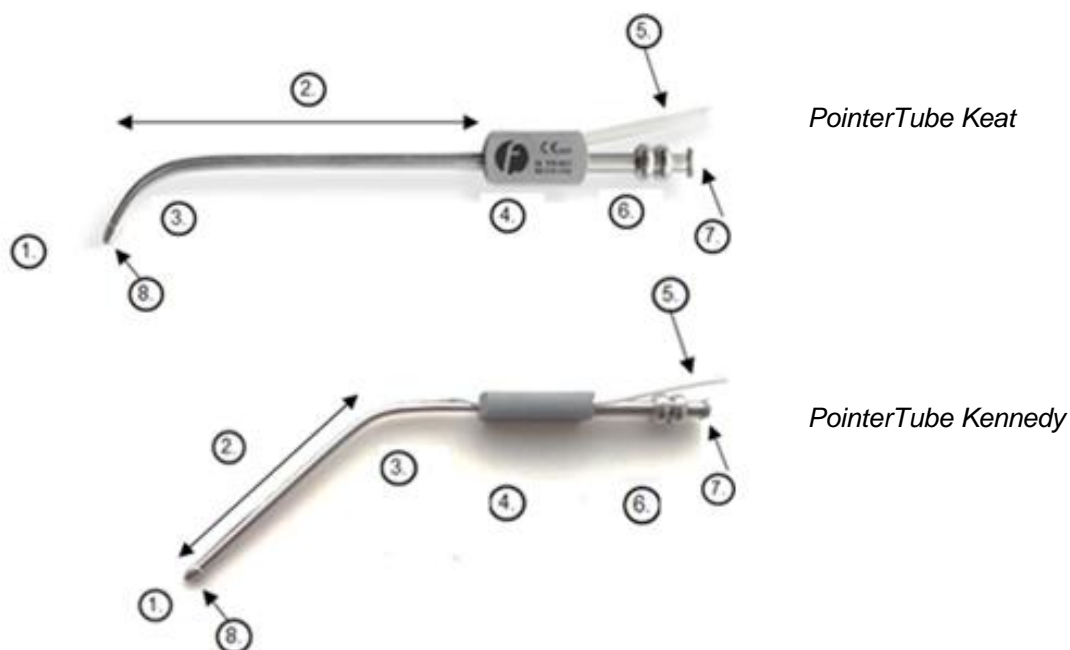
The instrument must not be exposed to MRI or used in a Magnetic Resonance Environment. The MRI exposure might magnetize the sensor. This might lead to a misleading navigation information.

3. Compatible devices

The instrument must only be used as a navigation instrument in connection with the navigation systems of Fiagon GmbH.

The instrument attaches to, and functions with, standard surgical vacuum suction systems. The maximum vacuum of the suction system should not exceed - 90 kPa.

4. Device Description



Description

1. Suction inlet
2. Tube length
3. Bending
4. Handle
5. Cable for connection with the navigation system
6. Flexible tube outlet (6-8 mm)
7. Suction outlet
8. Olive tip



CAUTION: The PointerTube Keat and the PointerTube Kennedy suction instruments cannot be bent.

5. Preparation for the navigation

After taking the instrument out of the sterile packaging, place it on the instrument table.

Connect the connector of the instrument with the navigation module. Note that the white marking at the plug aligns with the marking at the socket.



CAUTION: Note that you are inserting the plug of the instrument into a non-sterile device.

Once the instrument is connected, you will hear a confirmation tone and the status display of the instrument is displayed in color. The no of remaining uses is displayed in the status indicator of the instrument on the navigation screen. The instrument is accepted and registered to the navigation system.



NOTE: Before using the navigation the patient registration has to be done according to the description of the application. The instrument is allowed to be used for the patient registration. Consult the instructions for use

NOTE: If the instrument does not register at the navigation system, it cannot be used for the navigation.

NOTE: For connecting the instrument to the navigation unit via a longer distance use the extension cable instrument (accessory to the navigation unit).

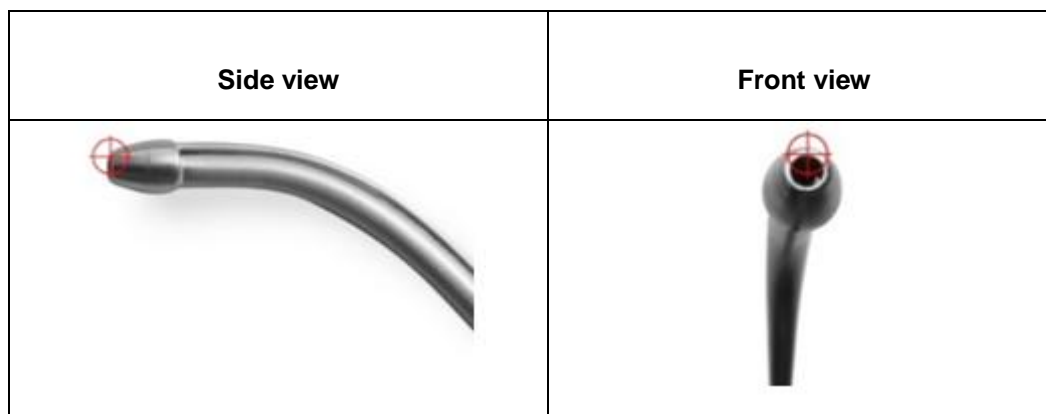
6. Use of the navigation functionality

After connecting the instrument to the system, the position of the navigation point is displayed on the navigation screen (if the patient registration has been performed before).



WARNING: Do not start using the navigation information of the instrument before you have checked and verified it. Therefore, check the displayed position of the instrument on several anatomical structures after connecting the instrument. If the deviation is significant do not use the instrument.

The figures below show the position of the navigation point on the suction instruments.



7. After the operation

After the operation disconnect the instrument from the navigation unit. Do this by pulling the plug directly. Do not pull the cable or the bend relieve. This might damage the cable.

Re-process the instrument according to the reprocessing instructions in these instructions for use.




CAUTION: The instrument should not be re-processes and reused after it has been used with patients with suspected Creutzfeldt-Jakob disease. There is a risk of disease transmission, dispose the instrument after the operation.

8. Specifications



PointerTubes

Type	PointerTube Kennedy E 01 2108	PointerTube Keat E 01 2107
Diameter outside	3 mm	3 mm
Diameter inside	2 mm	2 mm
Tube length	130 mm with olive tip	100 mm with olive tip
Suction outlet	6 - 8 mm	
Bendability	Not bendable	
Compatible suction machines	Suction machine ATMOS Record 500	
Operating conditions	Temperature: +15°C to + 35°C, Humidity: 30% to 75% without condensation Air pressure: 700 hPa to 1060 hPa	
Transport conditions	Temperature: -10°C to + 50°C, Humidity: 10% to 90% without condensation Air pressure: 700 hPa to 1060 hPa	
Storage conditions	Protected against dust, moisture and recontamination. Storage temperature: -10° to +50°C Storage humidity: 10% to 90% without condensation Storage air pressure: 700 hPa to 1060 hPa	

 To ensure that the instrument is intact, pay attention to the notes regarding the transport and storage conditions as well as the notes regarding the application of the instrument!

9. Reprocessing

Reprocessing instructions for reusable medical devices

<p>General information about the products</p> 	<p>The instruments are intended to be used with the Cube Navigation System. The usage of the products is described in the instructions for use of the Cube Navigation System.</p> <p>The instruments can be reprocessed and used 10 times. (see below for details)</p>
<p>Warnings</p> 	<p>Not qualified methods for cleaning and sterilization can damage cables and instruments.</p> <p>Instruments are supplied unsterile and must therefore undergo the complete reprocessing cycle prior to the first use.</p> <p>Pay attention to the notes and user manual of the sterilizer's manufacturer as well as the chemicals used.</p>
<p>Restrictions on Reprocessing</p>	<p>Sort out instruments that have been reprocessed 10 times.</p> <p>These instructions for use come with a track chart (count sheet) to keep record of the numbers of reprocessing cycles of an instrument. Each instrument has an individual serial number, which is labeled on the handle or the shaft.</p>

Instructions:

Site of use:	Remove surface soiling with a non-shedding disposable towel/paper towel. Do not allow saline, blood, body fluids, bone fragments or other organic debris to dry on instruments.
Storage and transport	Immediately after use on a patient, immerse the instrument in a cleaning agent/disinfectant (alkaline, free of aldehydes). Immersion the instrument prevents residues from drying (protein fixation). It is recommended to reprocess the instrument within 1h after use.
Preparation for reprocessing	The instruments are reprocessed as a complete unit with cable and connector. Cable or connector must not be removed.
Sorting out after 10 reprocessing cycles	Sort out instruments that have been reprocessed 10 times. To do so, these instructions for use come with a track chart (count sheet) to keep record of the numbers of reprocessing cycles of an instrument. Each instrument has an individual serial number, which is labeled on the handle or the shaft. When processing an instrument for the first time (brand-new), note the serial number on the first column of the track chart and check the tick box for the first reprocessing cycle. When reprocessing the same instrument, the next time, check the next tick box. When receiving an instrument for reprocessing that has a fully checked track chart, it has been reprocessed 10 times. Do not reprocess the instrument. Sort out the instrument and dispose it. (see section "Disposal" for details) Hint: You can use one track chart for ten instruments.
Cleaning - automated	Cannulated instruments "PointerTube" Equipment: <ul style="list-style-type: none"> · Washer/Disinfector equipped with instrument holders for cannulated instruments. (flexible tube connectors to Suction/Irrigation ports) · Washer type with following cycle: 5 minutes, 131°F (55°C), Injection rate, 0.64 fl.oz/gal (5 ml/l) of alkaline agent Deionized Water supply · Recommendation: AMSCO® 3052 Single-Chamber Washer/Disinfector (Steris, Inc.) · Cleaning agent: Neodisher® Mediclean forte · Deionized Water · Tap water, potable water Method: <ol style="list-style-type: none"> 1. Rinse each instrument thoroughly under warm (approximately 91°F/ 33°C) running tap water. Using a 10cc syringe, flush the ports of the cannulated instruments with tap water until there is no noticeable debris or discolored fluid exiting the instrument. 2. Place instruments in the instrument holders and connect the suction ports to the flexible tubes of the holder 3. Choose the instrument cycle in the washer. Validated: Wash: 5 minutes, 131°F (55°C), deionized water, Injection rate, 0.64 fl.oz/gal (5 ml/l) of alkaline agent (Prewash can be done additionally before the wash cycle) 4. Thermal Rinse at 194°F (90°C) for 3 minutes, deionized water

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Drying	Instruments can be dried until no visible moisture is present using filtered forced air for the inner lumen and using a clean, soft cloth for the outside of the instrument and the cable.
Cleaning inspection	Inspect the instrument for visible soil or debris. Repeat the cleaning steps in case residual soil or debris was found.
Inspection	The instrument should be inspected after cleaning and before use for surgery. Inspect the instrument visually for damage, wear and corrosion. Check also cable and connectors for damage. Sort out instruments that: <ul style="list-style-type: none"> • show indications of corrosion • have damaged cables or connectors • show tube denting Instruments that have been sorted out prior to their 10-cycles life time must not be used for surgery. These instruments can be disposed or returned to the dealer/manufacturer for the clarification of reason of the damage.
Maintenance	No special maintenance tasks are necessary for the instrument.
Repair, Returning instruments to the dealer/ manufacturer	Instruments are not being repaired during their life time. However, if you have sorted out an instrument during the inspection, inform the local dealer about the defect. The instrument might need to be returned to the dealer/manufacturer for the clarification of reason of the damage. Note that instruments must undergo the automated cleaning step before returning them.
Disposal	Used instruments must be disposed of as hazardous waste. Take steps to avoid risk of injury and infection. Protect against unauthorized access.
Packaging	Double pack each instrument in see-through peel pouches. Do not pack more than one instrument in a pouch. Recommended accessories: Pouches: Steriking (Healthmark Ind Co.) Size 7" x 12.5" and 8" x 15.75" Pouch item no.: #33 and #10 Bioindicator: MesaStrip Steam biological indicator (SGM Biotech, Inc.) Steriking is registered as a 510K device (K953776) with the FDA MesaStrip is registered with the FDA
Sterilization	Method: Prevacuum steam sterilizer Sterilization parameters: Min. temperature 270°F (132°C) Full cycle time: 4 minutes Min drying time 20 minutes Recommended validated Type: Middle-sized hospital sterilizer compliant to AAMI ST8 Size: 9 STE (sterilization units) Validated load mix: 2 container of instruments and 2 packages of linen
Storage	Make sure that the package is intact prior to storing. Store the instrument protected from dust, moisture and recontamination. The integrity of the sterile barrier needs to be protected.

Track chart (for 10 instruments)

Keep record of the number of reprocessing cycles.

Sort out instruments that have been reprocessed 10 times.

Use this track chart (count sheet) to keep record of the numbers of reprocessing cycles of an instrument. Each instrument has an individual serial number, which is labeled on the handle or the shaft.

When processing an instrument for the first time (brand-new), note the serial number on the first column of the track chart and check the tick box for the first reprocessing cycle.

When reprocessing the same instrument the next time, check the next tick box.

When receiving an instrument for reprocessing that has a fully checked track chart, it has been reprocessed 10 times. Do not reprocess the instrument.

Sort out the instrument and dispose it. (see section "Disposal" in the reprocessing instructions for details)

Hint: You can use one track chart for ten instruments.

Instrument SN no. Note here	Counter										
	Tick a box for each time the instrument with the SN number noted on the left is entering reprocessing										
	1	2	3	4	5	6	7	8	9	10	Dispose the instrument when all boxes of the line are ticked
	1	2	3	4	5	6	7	8	9	10	Dispose the instrument when all boxes of the line are ticked
	1	2	3	4	5	6	7	8	9	10	Dispose the instrument when all boxes of the line are ticked
	1	2	3	4	5	6	7	8	9	10	Dispose the instrument when all boxes of the line are ticked
	1	2	3	4	5	6	7	8	9	10	Dispose the instrument when all boxes of the line are ticked
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