

Glider Amplatz Sheath

Description:

Glider Amplatz Sheath

Configuration:

See brochure

Intended use:

Used for maintaining previously established nephrostomy tract.

Contraindications:

There are no known contraindications.

Complications:

- Tissue Trauma
- Tissue perforation
- Acute Bleeding
- Injury to the kidney

Caution:

Federal (USA) law restricts this device to sale by or on the order of a physician.

Warning/ Precaution:

- For single use only.
- Attempt to resterilize a single use device may compromise its structural integrity and/or lead to device failure which may result in patient injury or illness.
- Do not use if package is opened or damaged
- Dispose in accordance with recognized medical practice and under observance of applicable law and regulation
- Only experienced surgeons should use this device
- Storage temperature between 5°C to 30°C.
- Check compatibility with dilator before use.

Instruction for Use:

- After dilating the tract with the dilator of chosen size, advance the Amplatz sheath over the dilator in slow but firm twisting motion, till it enters into the renal pelvis.
- Once the position is confirmed via fluoroscopy remove the dilators leaving the guidewire in place, for establishing working tract.

Do not reuse, reprocess & resterilize:

Reuse, Reprocessing & resterilization may compromise the structural integrity, can also create risk of contamination & may not give desired result or create complications, infections which may result in injury, illness or death.

Limited Express Warranty:

The limited express warranty as set forth herein is exclusive and in lieu of all warranties of merchantability and fitness for use, remedies, obligations and liabilities for consequential damages. The products are being sold only for the purpose described herein and such limited express warranty runs only to the original user. In no event shall ALLWIN be liable for any breach of warranty in any amount exceeding the purchase price of the product. ALLWIN reserves the right to make design changes to products without liability to incorporate said changes in ALLWIN products previously designed or sold.



Catalogue
Number



Batch
Code



Date of
Manufacture



Use
By



Do not
re-use



Do not
Re-Sterilize

R_x only

Caution : Federal (USA) law
restricts device to sale by or
on the order of a physician.



Consult
Instruction
for use



Do not use
if Packing
is damaged



Caution



Keep out
of sunlight



Keep
Dry

STERILE EO

Sterilized using
ethylene oxide



30°C
Temperature
Limit



allwin
Medical Devices

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