Urinary Diversion Stents [Mono J]

Description:
Urinary Diversion Stent [Mono J]

Configuration:
See brochure

Intended use:
The Urinary Diversion Stent [Mono-J] is used for intraoperative placement to stent the ureter during percutaneous, endoscopy or operative procedures.

Contraindications:
Should not be used when it is too large to be advanced easily into the ureter.

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

Warning/ Precaution:

- Periodic evaluation is recommended. Not intended for permanent use.
- Check compatibility of the guidewire before use.
- Only experienced surgeon should place this device.
- Do not use if package is opened or damaged.
- Dispose in accordance with recognized medical practice and under observance of applicable law and regulation.
- 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.
- Storage temperature between 5ºC to 30ºC

Instruction for Use:
A variety of techniques may be used during stent placement. Each surgeon should use the methods with which user is most familiar.

For closed tip:
The stiff end of the guidewire is back-loaded onto the stent and the whole assembly is then advanced for appropriate placement.

For open tip:
The stent is advanced over the guidewire already placed.

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.

REF Catalogue Number | LOT Batch Code | Date of Manufacture | Use By | Do not re-use | Do not Re-Sterilize | Rx only — Caution: Federal Law restricts this device to sale by or on the order of a physician or a practitioner trained in its use.

Consult Instruction for use | Do not use if Packing is damaged | Caution | Keep out of sunlight | Keep Dry | STERILE EO — Sterilized using ethylene oxide

EC REP

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Temperature Limit

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