dispo foleys

INSTRUCTIONS FOR USE

Foley Balloon Catheter

MATERIAL USED :

- - PP.,SILICONIZED LATEX and PVC film

INDICATIONS :

- A Foley catheter is a thin, sterile tube inserted into the bladder to drain urine. It is held in place with a balloon at the end, which is filled with sterile water to prevent the catheter from being removed from the bladder. The urine drains through the catheter tube into a bag, which is emptied when full.

CONTRAINDICATIONS:

- Product should not to be used in patients with known Hypersensitivity to any of the materials used.
- Device must not be used other than as indicated in intended use.

INSTRUCTION FOR USE :

- Check the package sealing integrity to ensure sterility.
- During this procedure follow an aseptic technique using sterile gloves.
- Apply an appropriate sterile lubricant as per local policy.
- Carefully insert catheter tip into bladder until tip and balloon are suitably located (normally indicated by urine flow).
- Using a syringe without needle, inflate balloon with sterile/water saline solution - slowly inflate the balloon with the recommended volume to use, marked on this pack and on the funnel of the catheter, and gently pull the catheter shaft backwards to ensure that the balloon is correctly "seated" at the neck of the bladder.
- To deflate the balloon, push a syringe without needle into the valve. Allow the pressure within the balloon to push the plunger back and fill the syringe with water. If you notice slow or no deflation, gently re-seat the syringe. If needed, use gentle aspiration to encourage deflation. If permitted by hospital protocol, the balloon can be deflated by cutting the inflation line between the valve and the drainage connector.

WARNING :

- Do not expose to heat or direct sunlight.
- The product is guaranteed sterile & non-pyrogenic, if the package has not been opened or damaged. Keep away from direct sunlight and external heat sources.
- Twisting the catheter due to patient's movements or other cause may result in fluid path blockage.
- The use of this product is restricted to a qualified doctor or a Paramedic
- The product should be used according to instruction.
- After use, this product may be a potential biohazard. Handle and dispose off in accordance with accepted medical practice and applicable local, state and country laws and regulations.

CAUTION :

- This product contains natural Rubber latex which may cause allergic reaction.
- To be sold only by or on the the order of a physician. For single use only.
- If reused, the impaired functionality and/ or contamination can lead to injury or serve.

MANUFACTURER DISCLAIMS ANY RESPONSIBILITY FOR POSSIBLE CONSEQUENCES RESULTING FROM IMPROPER USE.

- The product should not be reprocessed.
- Visually inspect and carefully check the product and packaging before use. Improper transport and handling may cause structural and/or functional damage to device or packaging.
- The product is guaranteed non-toxic, sterile and non-pyrogenic if the packages is not opened or damaged.
- Do not clean or re-sterile. For single use only. Discard after use.
- The product should be used immediately after opening the packaging.
- The device must not be re-used after its usage, it can be a potential biohazard.
- Improper use can result in serious or fatal illness or injury.

STORAGE CONDITION :

- temperature 10°c to 45°c.

CE 0123

REF	Reference No.
MD	Medical Device
LOT	Lot No./Batch No.
	Manufacturer
	Date of Manufacturing
	Use By / Expiry Date
STERILE EO	Sterilised By Ethylene Oxide Single Sterile Barrier System
	Do Not Reuse
	Do Not Resterilize
	Non-Pyrogenic
	Do Not Use If Package is Damage
LATEX	This Product Contains Latex
	Caution
	Consult Instructions For Use
10°C 45°C	Temperature
EC REP	Authorised Representative In The European Community

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