INSTRUCTIONS FOR USE

PERITONEAL DIALYSIS TRANSFUSION SET

Materials Used

- Polypropylene, Acrylonitrile Butadiene Styrene (ABS), PVC, Latex Rubber, Stainless Steel needle.

Indications

- For purification of blood using the artificial kidney apparatus via filtration through an artificial semi permeable membrane.
- For removal of waste solution replaced by fresh solution.

Contraindications

- Not to be used in patients with known hypersensitivity to any of the materials used.

Directions for Use

- Close the regulating clamp.
- Remove the protection cap from the spike and insert the spike into the container to its full length.
- Squeeze the drip chamber and adjust the fluid level.
- Open the airvent caps of both spikes.
- Open regulating clamp. Expel the air from the tube and close regulating clamp.
- Connect the set with dialysis catheter and regulate the flow by adjusting regulating clamp to achieve desired flow rate.

Cautions

- Do not use if protective cap is loose or missing.
- Do not use if package is opened or damaged.
- Use Rubber part only for injecting the medicines.
- Close the airvent during periods of interrupted infusion therapy.
- The product should be used only by qualified healthcare professionals.
- Do not re-sterilize. Discard the set after single use.
- Reuse and cleaning of product may alter their structural and mechanical properties. It may lead to infection or other illness/injury.
- The product should be replaced and disposed of as per facility approved protocol or CDC guidelines.
- Store in between 5°C to 35°C, avoid excessive heat, protect from direct sunlight and moisture.
- Poly Medicure Limited will not be responsible for any direct incidental or consequential damages resulting from reuse of product.

Cautions

See Instructions for use

REF — Product reference/Art. No.

For single use only

Do not Resterilize

Do not use if packaging or product has been damaged or contaminated

LOT — Batch Number

Date of Manufacturing

Use by / Expiry date

STERILE EO — Sterilised by Ethylene Oxide

Single Sterile Barrier System

EU REP — European Authorised Representative

Manufactured by

Non Pyrogenic

5°C 35°C — Storage Condition

Keep Away from Sunlight

Keep Dry

MD — Medical Devices

POLYMED

EU REP Obelis S.a
Boulevard Général Wahis 53
1030 Brussels, BELGIUM
Tel : +(32) 2. 732.59.54
E-mail : mail@obelis.net

CE 0123

Not made with DEHP plasticizers

Rev. 06-25/08