dispoflow

I.V. Set with Flow Regulator Set

INSTRUCTIONS FOR USE

MATERIAL USED :

- PVC, ABS, PP, Latex, Silicon, Isoprene, Nylon mesh and SS

INDICATIONS :

- Infusion of I.V. Solutions.
- Intermittent intravenous drug administration under gravity feed through a needle or catheter inserted in vein.
- To maintain hydration and/or correct dehydration if patient is unable to take sufficient volume of oral fluid.

CONTRAINDICATIONS:

- Product should not to be used in patients with known Hypersensitivity to any of the materials used.
- Administration of high viscous fluids.
- Device should not be used for administration of blood and blood components.
- Device must not be used other than as indicated in intended use.

INSTRUCTION FOR USE :

- Check the package sealing integrity to ensure sterility.
- Remove the Infusion Set using aseptic technique.
- Open protective cover on the spike (closure piercing device).
- Insert spike into the out let of the solution container.
- Fill the drip chamber up to half by squeezing and releasing. Do not fill completely.
- Make sure that flow regulator is in 'on' position.
- Release clamp on IV set to allow fluid to flow through.
- Prime flow regulator with 10-20 ml.
- Re-clamp proximal (input) tube.
- Connect output tube to IV access.
- Set desired flow rate by rotating graduated marker and also by drop counting in drip chamber of I.V. Set and please note the drop size from the packing of I.V. Set flow rate have a tolerance of ±10%.
- Release clamp to allow fluid flow.
- Observe for adequate flow.

WARNING :

- 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.
- If there is any change in expected performance of the device or in case of any malfunction the device should be immediately removed & sent back to supplier for analysis

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 :

- Do not expose to heat or direct sunlight.

CE 0123

REF	Reference No.
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 & Consult Instructions For Use
10°C 45°C	Temperature Limitation
	Caution
	Consult Instructions For Use
EC REP	Authorised Representative In The European Community
MD	Medical Device

Disposafe Health And Lifecare Limited

Plot No.1 & 2, Phase-II, Sector-59, Ballabhgarh, Faridabad-121004, Haryana (INDIA)
E-mail : info@disposafe.com
Web : www.disposafe.com
Customer Care No. 91-129-4170201
Mfg. Lic. No. : MFG/MD/2019/000106

EC REP CEpartner4U BV., ESDOORNLAAN 13, 3951 DB MAARN, THE NETHERLANDS.
E-mail : office@cepartner4u.com

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