# INSTRUCTIONS FOR USE

## SCALP VEIN SET with/without Safety Features

**Materials Used**

- Polypropylene, Poly Vinyl Chloride (PVC), ABS, Polyethylene, Stainless Steel.

**Indications**

- Infusion of I.V. fluids and administration of other drugs and medicines.
- Maintaining hydration and/or correct dehydration in patients who are unable to take sufficient volume of oral fluids.

**Contraindications**

- Administration of highly viscous fluids.
- Not to be used in patients with known hypersensitivity to any of the materials used.

**Directions for Use**

1. Open the package.
2. Connect the funnel connector of scalp vein set to the male fitting of infusion set.
3. Grasp wings and remove the protective cap of needle of scalp vein set and open regulating clamp of I.V. set.
4. Expel the air from the PVC tube and close regulating clamp.
5. Perform venipuncture and confirm proper positioning in the vessel and regulate the flow by adjusting regulate the flow by adjusting regulating clamp to achieve desired flow rate.
6. After use, slide the safety cover towards needle until an audible click is heard and the safety feature is visually confirmed to be activated.
7. Do not attempt to deactivate the safety device by separating the needle from the safety cover.

**Note : Point no. 6 and 7 is applicable for Scalp Vein Set with safety feature.**

**Cautions**

- Do not use if package is opened or damaged.
- Check the product & package integrity prior use.
- Care should be taken not to touch the needle.
- Needle must be carefully and firmly placed into the lock position of the safety cover.
- Keep hands behind the needle at all times during use and disposal.
- The product should be used only by qualified healthcare professionals.
- Do not re-sterilize. Discard the set after single use.
- Some organic solvents, alcoholic disinfectants, infusion solutions and high pH-value substances might cause cracking of the female luer. It is recommended to follow the IFU of such medicine / disinfectant before using with the device.

  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.


POLYMED

**POLY MEDICURE LTD.**
Plot No.104-105, Sector -59, HSIIDC, Industrial Area,
Ballabhgarh Faridabad, Haryana- 121004,
INDIA, Web : www.polymedicure.com,
Contact customer care executive: +91-129-4287053,
email:customercare@polymedicure.com,
Address: Same as above

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

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

Single Sterile Barrier System

STERILE EO — Sterilised by Ethylene Oxide

EU REP — European Authorised Representative

Manufactured by

Non Pyrogenic

5°C 35°C — Storage Condition

Keep Away from Sunlight

Keep Dry

MD — Medical Devices

Not made with natural rubber latex
Not made with DEHP plasticizers

Rev. 00-25/08