Size = 158X120 mm, Black Print

INSTRUCTIONS FOR USE

Viggo Connect™
Extension Tube (Pressure Monitoring Tube (M/F))

MATERIALS USED:
• ABS, PP, PVC, HDPE, Polycarbonate (PC)

INDICATIONS:
• To add extra tube length in cases where IV set tubing is insufficient
• To separate the injection site from infusion site and prevent mechanical irritation of IV access (in patients at high risk for thrombophlebitis)
• To administer IV contrast agents/fluids in patients undergoing CT/MRI/PET scans, or other procedures that require patient to enter a gantry

CONTRAINDICATIONS:
• Product should not be used in patients with known hypersensitivity to any of the materials used

INSTRUCTIONS FOR USE:
•The connecting port of this device is in compliance to EN 20594-1 & ISO 594-2
• Attachment Of Extension Tube To IV Access
• Ensure that venous access (e.g. IV cannula) is patent by flushing with saline
• Inspect packing of Extension Tube visually to ascertain that package is intact
• Remove the tube from sterile packing taking care to maintain sterility
• Connect IV infusion line to the female luer; discard white injection stopper

Priming:
• Open clamps of IV infusion line and displace air entirely through ENTIRE LENGTH of EXTENSION TUBE. If required the priming can be done with normal saline separately to prevent wastage of infusion fluid
• Clamp the input IV line to stop flow through extension tube
• Remove protector from male luer at the distal end of extension tube
• Connect extension tube to venous access device (e.g. IV cannula) and secure the connection by rotating male luer lock
• Open clamp on IV line to allow flow of infusion fluid; observe for adequate flow

REMOVAL of EXTENSION TUBE
• Clamp input IV infusion lines to stop flow of fluid.
• Keep injection stopper ready for closing IV access (e.g. IV cannula) after removal
• Unscrew the male luer lock and remove extension tube; close IV access with injection stopper or other device
• Disconnect the female luer from input IV line

TERM OF USE :
• Recommended Maximum Duration of Use: Depends on type of medication and other factors; refer to your facility's guidelines or CDC Guidelines

WARNINGS
• The use of this product is restricted to a qualified doctor or a paramedic
• Read instructions before use
• The product should be used according to the instructions for use
• **For known/reported adverse events associated with use of this device, refer to the Clinical Evaluation Report HH-QA-CER-IV.**

VIGGO MEDICAL DEVICES LIMITED DISCLAIMS ANY RESPONSIBILITY FOR POSSIBLE CONSEQUENCES RESULTING FROM IMPROPER USE
• Any device that is connected to this product must comply with EN 20594-1 & ISO 594-2 in order to achieve the intended performance of this product & to avoid leakage in this connection
• 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 sterile & non-pyrogenic if the package has not been opened or damaged
• Do not clean or resterilise. FOR SINGLE USE ONLY. Discard after use
• Re-use of this device may change its mechanical or biological features and may cause device failure, allergic reactions or infections
• Contains phthalates, should not be used for the treatment of children, pregnant women or nursing mothers
• **Disposal/Discard:** Dispose off/Discard the used Device in accordance with your Country's Healthcare and Safety Regulations

VIGGO™
MEDICAL DEVICES

REF	Product Ref. No.
LOT	Batch Number
	Date Of Manufacturing
	Use By
2	Do Not Reuse
STERILE EO	Sterilised by Ethylene Oxide Gas
PHT DEHP	Contains Phthalates
	Caution, Consult Accompanying Documents
	Consult Instructions For Use
	Non-Pyrogenic
	Do Not Use if Package is Damage
2 STERILIZE	Do not resterilize
	Keep away from sunlight

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AW/IFU_ALa, Rev. 01
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