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INSTRUCTIONS FOR DISPOSABLE INFUSION SETS

INDICATIONS FOR USE

The product is gravity feed only and used to infusion.

Main structure and properties

-Tubing and drip chamber are transparent, can easy to observe;
-Tubing is flexible enough to resist winding and twisting;
-The design of closure-piercing device to make it convenient for handle and usage;
-Flow regulator can make the flow rate change from minimum to maximum.

USAGE

-Firstly, check valid period and completeness of package before using;
-Tear the package at the cut part and take out the infusion set;
-Close the regulator, pull out protective cap of closure-piercing device and push the spike into the disinfection closure of fluid container;
-Put fluid container upside down, press chamber and half of it was filled with injection solution; then open the regulator to expel air in tube.
-Penetrate vein according aseptic manipulation after expelling air then fix it.
-Observing and then change flow rate to normal.
-Infusion operation must have the professionals nursing to implement and guardian.
-20 drops=(1±0.1)ml[(1±0.1)g]

PRECAUTIONS

-It may causes hypothermia-esp, weak patients;
-It may causes abnormal thythem of ventrickes of heart because of fast flow rate;
-There should maintain the guardianship during the infusion in case of any accident;
-It should be sterile and free from pyrogen before package opened;.
-It has a valid period of five years, do not use if it is beyond valid date.
-Discard after singe use.

WARNINGS

-Do not use if the package is opened or damaged, or protective cap is aparted, any part is missed, and foreign matter is found;
-It should be used by special trained person, and the skin should be correctly sterilized.
-The products can not be reused. The repetition use may cause cross infection, even serious illnesses such as hepatitis, HIV and other infectious disease.
-Always dispose of blood contaminated products in a manner consistent with established biohazard procedures.

STORAGE

Product should be kept in a condition that relative humidity is less than 80%, it also should be free from corrosive gas, cool, dry with good ventilation and clean.

Any serious incident that has occurred in relation to the device should be reported to the manufacturer and the competent authority of the Member State in which the user and/or patient is established.

UDI Unique device identifier	LOT Batch code	REF Catalogue number	Caution	STERILE EO Sterilized using ethylene oxide	
Date of manufacture	Manufacturer	Consult Instructions For Use	Do not re-use	Do not use if package is damaged	
MD Medical device	Use-by date	Do not resterilize	Fragile, handle with care	Temperature limit	CE 2862
Keep away from sunlight	Keep dry	Non-pyrogenic	This way up	Humidity limitation	
Single sterile barrier system	STERILE Sterile	PHT DEHP Contains or presence of Phthalate		REV.: C/0 EFFECTIVE DATE: 2025-05-30	

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