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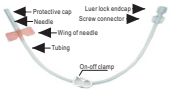
# Tkmd® Instructions for A.V. Fistula Needles

**【Scope of Application】**

・This product is suitable for establishing vascular access in hemodialysis treatment.

**【Main structure and properties】**

・The procduct is made up of protective cap, needle stand, tubing, on-off clamp, luer lock connector And luer lock endcap;
・This product has a good property of rigidity, toughness and corrosion resistance;
・Tubing is transparent that can easy to observe;
・Tubing is flexible enough to resist winding and twisting;
・On-off clamp can make the blood rate change from minimum to maximum;
・The design of needle stand to make it convenient for handle.




**【Specification】**

・15G～17G

**【Usage】**

・ Firstly, check expiry period and completeness of the products before using;
・Open the package from the peel site and take out A.V. fistula Needles;
・Connect connector with pipe of machine;
・Pull out needle protective cap and expel air in tubing;
・Push the needle spike into the body vein according aseptic manipulation and fix it.

**【Precautions】**

・It may causes hypothermia especially to weak patients;
・It may causes abnormal rhythm of ventricles of heart because of fast flow rate;
・There should be guardian during 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 and Indications】**

・Do not use if the package has been opened or damaged, protective cap has been removed, there are missing parts, and if foreign matter is present.
・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.

**【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.

**【Symbols】**

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|---|---|---|---|---|---|
| UDI Unique device identifier | LOT Batch code | REF Catalogue number | Contains or presence of phthalates | STERILE EO Sterilized using ethylene oxide | |
| Date of manufacture | Manufacturer | Do not re-use | Caution | 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 | | REV.: B/0 DATE: 2025-05-30 | |


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