



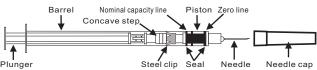
INSTRUCTIONS FOR Auto Disable Syringes

INDICATIONS FOR USE

The auto disable syringe is intended for use in the suction and injection of vaccine for medical purposes. Additionally, after injection to the body, the plunger can be automatically locked by the mechanism to prevent the re-use of this syringe.

Patient target group or groups: All population.

MAIN STRUCTURE, PERFORMANCE CHARACTERISTICS AND EXPECTED CLINICAL BENEFITS



-Barrel is transparent, smooth and can be observed liquid clearly.

-In the same plane, the plunger can be visible observed same-distance concave step, which set destructive area.

-Steel clip, between the barrel and plunger, is destructive after using.

-The syringe itself has fixed connected hypodermic needle.

-Expected clinical benefits: on-demand smart and precise drug delivery; reduced systemic side effects like redness, pain, swelling; reduced risk of needle stick injuries.

-Performance characteristics:Precision dose injection,Avoid secondary use and cross contamination. **Specification:** TKADS01: 0.05ml. Needle size: 23G to 31G.

USAGE

 Peel open sterile wrapper.
Remove the protective cap without touching the needle.
Do not pull plunger backward before using the syringe. 4) Insert needle in clean rubber cap of vaccine vial and draw up a dose.
Expel air or excee vaccine, adjusting plunger to the

indicated vaccine dose.

6) Inject vaccine.

7) Pull the syringe back.8) Place the used syringe into the safety box (do not ove-fill the safety box)

PRECAUTIONS

-The needle upwardly and expel all the air in syringe barrel, in case air be injected into human body.

-Abnormal operation, do not pull plunger. Sucking liquid, do not push back. because it will make the product destroyed. -It should be sterile and non-pyrogenic before package opened.

-It has a valid period of five years. Do not use if it is beyond valid date.

WARNINGS

-Do not use if the package is opened or damaged, or protective cap is aparted or 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.

-A notice to the user and/or patient that 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.

STORAGE

-Product should be kept in a condition that relative humidity is less than 80%, temperature ranges from -10 to 40 $^\circ\!C.$ SYMBOLS

UDI Unique device identifier	LOT Batch code	REF Catalogue number	Caution	STERILEEO Sterilized using ethylene oxide	
쎄 Date of manufacture	Manufacturer	(1) Re-use prevention	🚫 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	
🔆 Keep away from sunlight	 ″ Keep dry	Non-pyrogenic	↑↑ This way up	Humidity limitation	L Z 2862
Unique sterile barrier system		Simple aseptic barrier system with external protective packaging		REV.: B/0 EFFECTIVE DATE: 2023-06-30	

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