

# Instruction for Use

## **【Production Name】**

Pressure Sensitive Tape

## **【Model】**

C150 \C151 \C152 \C153 \C154 \C155 \C156

## **【Product feature, Main Structural Composition/Constituents】**

Tape usually coated with self-adhesive properties on the back material. Part of the adhesive tape coated with a protective layer. Nonsterile supply, single use. No direct contact with the wound. The sticking site is good skin.

## **【Range of Application】**

It is used for attaching dressings to wounds or fixing other medical instruments to specific parts of the human body.

## **【Cautions】**

1. It is prohibited for skin allergic to pressure sensitive glue.
2. This product is a foreign product, no entry.
3. If it is necessary to increase viscosity at low temperature, it can be warmed slightly.
4. Children should use it under the guidance and supervision of their parents.
- 5.. This product is disposable.
6. Please dispose of the product as medical waste after use.

## **【product maintenance and maintenance method, special storage, transportation conditions and methods】**

Store in a closed place with dry, non-corrosive gas at normal temperature. Do not take the product out of the bag or box and store it naked in the air. Use vehicles to transport at room temperature, prevent rain, forbid direct sunlight.

**【Production Address】**

No.1 Guangdong Road, Jiangshan Industrial Park, Laixi, Qingdao, China, 266603

**【Production Enterprise】**

Qingdao Hainuo Biological Engineering Co., Ltd.

**【Production Date / Batch Number, Service Life】**

Refer to the Package.









**【Product Validity】**







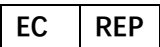
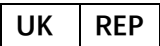
3 Years.

**【Sterilization】**

Non-sterile.

**【Symbol Description】**

	Caution, Indicates the need for the user to consult the instructions for use for important cautionary information, such as warnings and precautions that cannot, for a variety of reasons, be presented on the medical device itself.
	CE Mark: conforms to essential requirements of the Medical Device Regulation 2017/745 (EU)
	UKCA Mark: conforms to essential requirements of the Medical Device Regulation Part II of the UK MDR 2002
	Date of manufacture, Indicates the date when the medical device was manufactured
	Manufacturer, Indicates the medical device manufacturer, as defined in Medical Device Regulation 2017/745 (EU)
	Use-by date, Indicates the date after which the medical device is not to be used
	Do not re-use, indicates a medical device that is intended for one use, or for use on a single patient during a single procedure
	Do not use if package is damaged, indicates a medical device that should not be used if the package has been damaged or opened

	Non-sterile
UDI-DI	UDI-DI code of product identification
	Batch code, Indicates the manufacturer's batch code so that the batch or lot can be identified.
	Keep away from sunlight, indicates a medical device that needs protection from light sources.
	Keep dry, indicates a medical device that needs to be protected from moisture.
	Medical Device mark
	<p>Qingdao Hainuo Biological Engineering Co., Ltd.</p> <p>No.1 Guangdong Road, Jiangshan Industrial Park, Laixi, Qingdao, China.</p>
	<p>CMC Medical Devices &amp; Drugs S.L</p> <p>C/Horacio Lengo N° 18 CP 29006, Málaga-Spain</p>
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