Instruction for Use

【Product Name】

Alcohol Prep Pad

【Model】

C204,C205,C206,C207,C208,C209

【Size】

6*3/6*6/12*12cm

【Intended use】

Alcohol Prep Pad is used to disinfect the skin, around the wound and before injection.

【Raw Material】

1) Spunlace non-woven fabric
2) Saturated with 70%(± 5%) Isopropyl alcohol

【Directions】

Wipe topically and discard after use

【Caution】

1) Do not use
   - With electrocautery procedures.
   - In the eyes
   - If contact occurs flush eyes with water
2) Stop use
   - If irritation and redness develop, and if symptoms last, pls consult your health care practitioner.
3) Keep out of reach of children
   - If swallowed, get medical help or contact a Poison Control Center right away.

【Warnings】

1) This product is for single use only
2) For external use only. Flammable, keep away from fire or flame.
3) If any adverse event occurs, it should be reported to the manufacturer and competent authority of the Member State where the user and/or patient is located.
4)

【Package】

The inner packaging uses paper bags, the medium package is a white cardboard and 50 pads/box

【Expiry】

Two years.

【Production Date】

Refer to the package label.

【Storage conditions】

Store at room temperature 15℃~30℃(59°~86°F)

【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
LOT	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.
MD	Medical Device mark
	Qingdao Hainuo Biological Engineering Co., Ltd. No.1 Guangdong Road, Jiangshan Industrial Park, Laixi, Qingdao, China.
EC REP	CMC Medical Devices & Drugs S.L C/Horacio Lengo Nº 18 CP 29006, Málaga-Spain
UK REP	Kingsmead Service Limited 19 Mezzanine Floor 19-21 Crawford Street London England W1H 1PJ