Instruction for Use

【Production Name】

Wound Plasters

【Model】

C101\C102\C103\C104\C106\C107\C108\C109\C110\C111\C112\C113\C114\C115\C116\C117

【Brief Introduction】

Wound plasters is mainly composed of flat cloth and absorbent pads. It is used for hemostasis and protecting creation.

【Intended Use】

It is used to block and isolate skin contact with the outside world, and bandaging nursing.

【Contraindications】

Do not apply it to open wounds on the skin. If you have skin allergies or feel uncomfortable, stop using it immediately.

【Sterilization】

Non-sterilization.

【Precautions, Warnings and Prompts】

1. The Wound Plasters is disposable product, it cannot be reused that preventing cross infection.
2. The packaging is prohibited to use if is damaged.
3. Do not use expired products.
4. Disinfect the wound before use, and do not touch the protective pad with your hands after unpacking.

【Instructions for Use】

1. According to the users' wound size, select the appropriate specifications of wound plasters.
2. After the package is opened, the small package can be removed.
3. Clean the wound surface, tear off the release paper of the product, stick the protective pad on the wound surface, fix the position of the adhesive backing material, and tighten appropriately.

【Storage Conditions】

Storage should not exceed 85% of the relative humidity, no corrosive gases and good ventilation indoors.

【Transport Conditions】

Transported by making use of means of transportation under normal temperature, to prevent rain.

【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.

【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