CE

Instruction for Use

【Product Name】

Disposable Nitrile Gloves

【Model】

C610（S,M,L,XL,XXL）

【Material】

Nitrile Butadiene Rubber

【Intended use】

It is intended to provide a barrier to help prevent cross-contamination of infectious agents between the visitor and patients.

【Contraindications】

People who are allergic to product materials.

【Directions for use】

- How to put on gloves

1. Select the appropriately sized gloves.
2. Check the validity period of the gloves on the package and the tightness of the package.
3. Wash hands or sanitize before wearing gloves.
4. Open the glove package. Lift the opening of the package with one hand, and take out the gloves with the spread fingers and insert hand into glove.
5. Lift the opening of anther package by the same way above. Hold another glove with the gloved hand and insert the other hand into the glove. Put your hands together, and adjust the wearing of gloves by crossing your fingers.

- How to remove the gloves

1. Grab the end of the outsaid of glove with gloved fingers and pull the glove down.
2. Hold the removed glove with the gloved hand, and grab the end of the inside of glove with the non-gloved hand to pull the glove down.
3. Hold the removed gloves inside out. Throw away the gloves in a medical waste

place and wash your hands.

【Warnings】

1) This product is disposable and destroyed after use.
2) It is forbidden to use if the package is damaged.
3) This product does not have flame retardant performance.
4) 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.

【Package】

The inner packaging uses plastic packaging bags, the medium package is a carton and 100 pcs per box.

【Expiry】

5 years

【Production Date】

Refer to the package label.

【Storage conditions】

The product shall be stored in a room without corrosive gas and with good ventilation.

【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