

## EC Declaration of Conformity

**Manufacturer:**

**Name:** Xianning Full Guard Medical Products Co., Ltd

**Address:** Yong'an East Avenue, Xian'an Economic Development Zone, Xianning City, Hubei Province, China

**Tel/Fax:** 0715-8200113

**SRN:** Not available yet

**Whose single Authorized Representative:**

**Name:** ZOUSTECH S.L

**Address:** Pso.Castellana, 141- planta 19, 28046-Madrid, Spain

**Tel/Fax:** +34694426446

**SRN:** ES-AR-000002008

### **Disposable Medical Hole Towel**

**UMDNS CODE:** 15646

**Product Code:** FDP

**Product Size:** 50x50cm, 75x90cm, 100x150cm, or customized size

**Classification According To MDD, Annex VII:** Class I Sterile, Rule 1

**Applied Common Specification/Standard:**

EN 13795-1:2019 Surgical clothing and drapes-Requirements and test methods part1:Surgical drapes and gown

EN ISO 11607-1:2020 Packaging for terminally sterilized medical devices - Part 1: Requirements for materials, sterile barrier systems and packaging systems

EN ISO 11607-2:2020 Packaging for terminally sterilized medical devices - Part 2: Validation requirements for forming, sealing and assembly processes

EN ISO 11135:2014 Sterilization of health-care products - Ethylene oxide - Requirements for the development, validation and routine control of a sterilization process for medical devices

EN ISO 14971:2019 Medical devices - Application of risk management to medical devices (ISO 14971:2019)

ISO 10993-1:2018 Biological evaluation of medical devices - Part 1: Evaluation and testing within a risk management process

EN ISO 10993-5:2009 Biological evaluation of medical devices - Part 5: Tests for in vitro cytotoxicity (ISO 10993-5:2009)

EN ISO 10993-10:2013 Biological evaluation of medical devices - Part 10: Tests for irritation and skin sensitization (ISO 10993-10:2010)





EN 1041:2008 Information supplied by the manufacturer of medical devices

EN ISO 15223-1:2016 Medical devices—Symbols to be used with medical device labels, labeling and information to be supplied.

We, the manufacturer, herewith declare under our sole responsibility that the above-mentioned product, meets the provision of the following EC Council Directives and All applicable harmonized Standards. All supporting documentations are retained under the premises of the manufacturer.

**DIRECTIVES**

Medical Device Directive:

COUNCIL DIRECTIVE 93/42/EEC of 14 June 1993 concerning medical devices (MDD 93/42/EEC),  
Amended by DIRECTIVE 2007/47/EC of 5 September 2007.



Notified Body: TÜV SÜD Product Service GmbH, GmbH, Ridlerstr: 65, 80339, München, Germany

NB Identification number: 0123

Certificate No.: G2S 003747 0002 Rev.00

(EC) Certificate(s): YES

Expire date of the Certificate: Nov 04, 2023

Start of CE Marking: Nov 05, 2018

Place of Issue: Xianning, Hubei 湖北咸宁

Date of Issue: 2021. 5.26

Signature:

Name: 姜迪 Rosen Jiang

Position: Managing Director

Stamp: