

## **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:** CN-MF-000013685

**Whose single Authorized Representative:**

**Name:** ZOUSTECH S.L

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

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**SRN:** ES-AR-000002008

### **Disposable Medium Drapes**

**UMDNS CODE:** 12368

**Product Code:** FGSD

**Product Size:** (30-300)×(30-400) cm

**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 ISO 1041:2008**    Information supplied by the manufacturer of medical devices

**EN ISO 15223-1:2016**    Medical devices – Symbols to used with medical devices 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 Sep 2007.



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

NB Identification number: 0123

Certificate No: G2S 003747 0002 Rev.00

(EC) Certificate(s): YES

Expire date of Certificate: Nov 04, 2023

Start of CE Marking: Nov 05, 2018

*Place of Issue: Xiannng, Hubei*

*Date of Issue: 2021.5.26*

*Signature:*

*Name: 姜迪 Rosen Jiang*

*Position: Managing Director*

*Stamp:*

