EU Declaration of Conformity

Manufacturer Name: Kingstar Medical (Xianning) Co., Ltd.
Manufacturer Address: No. 79 Yong'andong Road, Xian'an District 437100, Xianning City, Hubei Province, the People's Republic of China
SRN of the Manufacturer: CN-MF-000006015
Location of Manufacturer: Xianning City, Hubei Province, China.
Authorized Representative: Shanghai International Holding Corp. GmbH (Europe) [DE]
SRN of the Authorized Representative: DE-AR-000000001
Address of their Registered Place of Business: Eiffestraße 80, 20537 Hamburg, Germany
Location be established: Germany

Basic UDI-DI: 693666261221001RZ
Trade Name: Sterile Absorbent Pad
Name of the device: Sterile Absorbent Pad
EMDN Code: M04010101, Non-Woven Adhesive Dressings, With Absorbent Pad
MDR Certificate No.: G11 097364 0014 Rev. 02

Intended Purpose: Intended to be covered on skin surface near wound to absorb blood and/or exudates during the treatment of the wound.

Risk Class of the Device: Class I sterile, based on Rule 4 of ANNEX VIII of Regulation (EU) 2017/745.
-All non-invasive devices which come into contact with injured skin or mucous membrane are classified as class I if they are intended to be used as a mechanical barrier, for compression or for absorption of exudates.

The conformity assessment procedure performed: Because the devices are placed on the market in sterile condition, the procedures set out in Chapters I and III of Annex IX are applied. The notified body involved to the aspects relating to establishing, securing and maintaining sterile conditions.

CS used or Standard applied: Please find in Annex II.
Identification of the device: Please find in Annex I.

Declaration: This declaration of conformity is issued under the sole responsibility of Kingstar Medical (Xianning) Co., Ltd. We hereby declare that the medical device specified above meet the provision of the Regulation (EU) 2017/745 for medical device. This declaration is supported by the quality system approval to ISO 13485 by TÜV SÜD Product Service GmbH.
All supporting documentation is retained at the premises of the manufacturer.

Notified Body: TÜV SÜD Product Service GmbH
Address: Ridlerstraße. 65, 80339 Munich, Germany
Identification No.: CE0123

Signed for and on behalf of:
Place of Issue: Xianning City, Hubei Province, China.
Date of Issue: 2026.03.24
Print Name: Fan Rong
Function: Management Representative
Signature: 

Annex I --- Identification of the Device Covered by the EU Declaration of Conformity

1. Identification of the Device

Table --- Identification of the Device

No.	REF No.	IVF Hartmann Art.	Specification	Packaging Configuration
1	C143020	2854200	3.5cm×5cm-12ply	2 pcs/pouch, 140 pouches/box, 16 boxes/carton
2	C143021	2853000	3.5cm×5cm-12ply	3 pcs/pouch, 120 pouches/box, 16 boxes/carton

Annex II --- European Harmonization and International Standard list

No.	Standards	Content
1	EN ISO 13485:2016/A11: 2021	Quality management systems — Requirements for regulatory purposes (ISO 13485:2016)
2	ISO 15223-1: 2021/Amd1: 2025	Medical devices - Symbols to be used with information to be supplied by the manufacturer- Part1: General requirements - Amendment 1: Addition of defined term for authorized representative and modified EC REP symbol to not be country or region specific
3	EN ISO 20417: 2021	Medical devices- Information to be supplied by the manufacturer (ISO 20417:2021)
4	EN ISO 14971:2019/A11: 2021	Medical devices-Application of risk management to medical devices (ISO 14971:2019)
5	EN 62366-1:2015/A1: 2020	Medical devices-Application of usability engineering to medical devices - Amendment 1
6	MEDDEV 2.7.1 rev 4	CLINICAL EVALUATION: A GUIDE FOR MANUFACTURERS AND NOTIFIED BODIES UNDER DIRECTIVES 93/42/EEC and 901385/EEC
7	ISO 2859-1:1999/A1: 2011	Sampling procedures for inspection by attributes--Part 1: Sampling schemes indexed by acceptance quality limit (AQL) for lot-by-lot inspection - Amendment 1
8	ISTA 2A: 2011	Partial simulation performance test procedure
9	MEDDEV 2.12-1: 2013	Guidelines on a medical devices vigilance system
10	ISO/TR 20416: 2020	Medical devices - Post-market surveillance for manufacturers
11	EN ISO 14644-1: 2015	Cleanrooms and associated controlled environments — Part 1: Classification of air cleanliness by particle concentration (ISO 14644-1:2015)
12	EN ISO 14644-2: 2015	Cleanrooms and associated controlled environments - Part 2: Monitoring to provide evidence of cleanroom performance related to air cleanliness by particle concentration (ISO 14644-2:2015)
13	ISO 14644-3:2019/Amd 1: 2020	Cleanrooms and associated controlled environments - Part 3: Test methods (ISO 14644-3:2019)
14	ISO 10993-1: 2025	Biological evaluation of medical devices Part 1: Requirements and general principles for the evaluation of biological safety within a risk management process
15	EN ISO 10993-5: 2009	Biological evaluation of medical devices - Part 5: Tests for in vitro cytotoxicity (ISO 10993-5:2009)
16	EN ISO 10993-7:2008/A1: 2022	Biological evaluation of medical devices - Part 7: Ethylene oxide sterilization residuals - Amendment 1: Applicability of allowable limits for neonates and infants (ISO 10993-7:2008/Amd 1:2019)
17	EN ISO 10993-10: 2023	Biological evaluation of medical devices - Part 10: Tests for skin sensitization (ISO 10993-10:2021)
18	EN ISO 10993-11: 2018	Biological evaluation of medical devices - Part 11: Tests for systemic toxicity (ISO 10993-11:2017)
19	ISO 10993-18:2020/A1: 2023	Biological evaluation of medical devices - Part 18: Chemical characterization of medical device materials within a risk management process - Amendment 1: Determination of the uncertainty factor (ISO 10993-18:2020/Amd 1:2022)
20	EN ISO 10993-23: 2021	Biological evaluation of medical devices - Part 23: Tests for irritation (ISO 10993-23:2021)
21	EN ISO 11135:2014/A1: 2019	Sterilization of health-care products - Ethylene oxide - Requirements for the development, validation and routine control of a sterilization process for medical devices - Amendment 1: Revisionof Annex E, Single batch release (ISO 11135:2014/Amd 1:2018)

22	EN ISO 11607-1:2020/A1: 2023	Packaging for terminally sterilized medical devices - Part 1: Requirements for materials, sterile barrier systems and packaging systems - Amendment 1: Application of risk management (ISO 11607-1:2019/Amd 1:2023)
23	EN ISO 11607-2:2020/A1: 2023	Packaging for terminally sterilized medical devices - Part 2: Validation requirements for forming, sealing and assembly processes - Amendment 1: Application of risk management (ISO 11607-2:2019/Amd 1:2023)
24	EN ISO 11737-1:2018/A1: 2021	Sterilization of health care products - Microbiological methods - Part 1: Determination of a population of microorganisms on products -Amendment 1 (ISO 11737-1-2018/Amd 1:2021)
25	EN ISO 11737-2: 2020	Sterilization of health care products - Microbiological methods - Part 2: Tests of sterility performed in the definition, validation and maintenance of a sterilization process (ISO 11737-2:2019)
26	EN 868-5: 2018	Packaging for terminally sterilized medical devices --- Part 5: Sealable pouches and reels of porous materials and plastic film construction - Requirements and test methods
27	EN ISO 11138-1: 2017	Sterilization of health care products --- Biological indicators --- Part 1: General requirements (ISO 11138-1:2017)
28	EN ISO 11138-2: 2017	Sterilization of health care products - Biological indicators - Part 2: Biological indicators for ethylene oxide sterilization processes (ISO 11138-2:2017)
29	REACH Regulation 1907/2006	Registration, Evaluation, Authorization and Restriction of Chemicals
30	CLP Regulation 1272/2008	Classification, Labelling and Packaging of Substances and Mixtures, Amending and Repealing Directives 67548/EEC and 1999/45/EC, and amending Regulation (EC) No 190712006
31	EN 1644-1: 1997	Test methods for nonwoven compresses for medical use - Part 1: Nonwovens used in the manufacture of compresses
32	EN 1644-2: 2000	Test methods for nonwoven compresses for medical use - Part 2: Compresses