

|              |                |             |                                                           |            |       |
|--------------|----------------|-------------|-----------------------------------------------------------|------------|-------|
| Doc. No.     | KSX/TD-ADS-017 | Title       | EU Declaration of Conformity of Sterile Adhesive Dressing |            |       |
| Ver./Rev.No. | A/0            | Issued Date | 2023.02.15                                                | Page/Total | 1 / 3 |

## 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)

**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:** 6971872201231001M6

**Name of the device:** Sterile Adhesive Dressing

**EMDN Code:** M0401, Prepared Dressings

**UMDNS Code:** 10276, Bandages/ Dressings, Adhesive

**GMDN Code:** 34864, Adhesive bandage

**Intended Purpose:** Sterile Adhesive Dressing is a device intended to be applied to a part of a patient's body and held in place by its pressure-sensitive adhesive to secure objects to the skin, cover and protect the skin edges of a 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:** Ridlerstr. 65, 80339 Munich, Germany

**Identification No.:** CE0123

**EC-Certificate No.:** G11 097364 0014 Rev. 00

**Certificate Valid from:** 2023-02-10

**Certificate Valid until:** 2028-02-09

**Signed for and on behalf of:**

Place of Issue: Xianning City, Hubei Province, China.

Date of Issue: 2023.02.15

Print Name: Fan Rong

Function: Management Representative

Signature: 

|              |                |             |                                                           |            |       |
|--------------|----------------|-------------|-----------------------------------------------------------|------------|-------|
| Doc. No.     | KSX/TD-ADS-017 | Title       | EU Declaration of Conformity of Sterile Adhesive Dressing |            |       |
| Ver./Rev.No. | A/0            | Issued Date | 2023.02.15                                                | Page/Total | 2 / 3 |

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

### 1. Identification of the Device

Table --- Identification of the Sterile Adhesive Dressing

| Classification | Meditrade Code | Meditrade Code            | Kingstar name                    | Kingstar REF |
|----------------|----------------|---------------------------|----------------------------------|--------------|
| Is             | 4457           | ABE Kanülenfixierung I.V. | Sterile Adhesive Dressing series | C133005      |
| Is             | 4752           | ABE Wundverband           | Sterile Adhesive Dressing series | C133007      |

### 2. Photograph of Sterile Adhesive Dressing

Drawings and photos of device:



Photo 1 ---Sterile Adhesive Dressing (type 1)  
2)



Photo 2 --- Sterile Adhesive Dressing (type 2)

## Annex II --- European Harmonization and International Standard list

| No. | Reference and title of the standard (and reference document)                                                                                                                                    | First publication<br>[1] | Reference of superseded standard |
|-----|-------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|--------------------------|----------------------------------|
| 1   | EN 556-1: 2001 Sterilization of medical devices - Requirements for medical devices to be designated 'STERILE' - Part 1                                                                          | 31.7.2002                | EN 556: 1994 + A1: 1998          |
| 2   | EN 556-1: 2001/AC: 2006                                                                                                                                                                         | 15.11.2006               |                                  |
| 3   | EN ISO 15223-1:2021 Medical devices — Symbols to be used with medical device labels, labelling and information to be supplied — Part 1: General requirements                                    | 06/07/2021               | EN ISO 15223-1:2016              |
| 4   | EN 1041:2008+A1:2013 Information supplied by the manufacturer of medical devices                                                                                                                | 25.9.2013                | EN 1041: 1998                    |
| 5   | EN 1422: 2014 Sterilizers for medical purposes - Ethylene oxide sterilizers - Requirements and test methods                                                                                     | 17.04.2014               | EN 1422: 1998                    |
| 6   | EN ISO 10993-1: 2018 Biological evaluation of medical devices - Part 1: Evaluation and testing within a risk management process.                                                                | 2018-08                  | EN ISO 10993-1: 2009             |
| 7   | EN ISO 10993-1: 2018/AC: 2010                                                                                                                                                                   | 18.1.2011                |                                  |
| 8   | ISO 10993-1: 2018 Biological evaluation of medical devices - Part 1                                                                                                                             | 08.2018                  | ISO 10993-1: 2009                |
| 9   | EN ISO 10993-5: 2009 Biological evaluation of medical devices - Part 5: Tests for in vitro cytotoxicity                                                                                         | 20/05/2009               | EN ISO 10993-5: 1999             |
| 10  | EN ISO 10993-10: 2013 Biological evaluation of medical devices Part 10: Tests for irritation and skin sensitization (ISO 10993-10:2010)                                                         | 21.8.2013                | EN ISO 10993-10: 2010            |
| 11  | EN ISO 10993-7: 2008 Biological evaluation of medical devices - Part 7: Ethylene oxide sterilization residuals                                                                                  | 2008-10                  | EN ISO 10993-7: 1995             |
| 12  | ISO 10993-7:2008/Amd 1:2019 Biological evaluation of medical devices — Part 7: Ethylene oxide sterilization residuals — Amendment 1: Applicability of allowable limits for neonates and infants | 2019-12                  |                                  |
| 13  | EN ISO 11138-2: 2017 Sterilization of health care products - Biological indicators - Part 2: Biological indicators (ISO 11138-2: 2017)                                                          | 29.3.2017                | EN ISO 11138-2: 2009             |

| Doc. No.     | KSX/TD-ADS-017 | Title       | EU Declaration of Conformity of Sterile Adhesive Dressing |            |       |
|--------------|----------------|-------------|-----------------------------------------------------------|------------|-------|
| Ver./Rev.No. | A/0            | Issued Date | 2023.02.15                                                | Page/Total | 3 / 3 |

|    |                                                                                                                                                                                                                                                  |            |                              |
|----|--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|------------|------------------------------|
| 14 | EN ISO 11140-1: 2014 Sterilization of health care products - Chemical indicators - Part 1: General requirements (ISO 11140-1: 2014)                                                                                                              | 12.11.2014 | EN ISO 11140-1: 2009         |
| 15 | EN ISO 11607-1:2020 Packaging for terminally sterilized medical devices - Part 1: (ISO 11607-1: 2019)                                                                                                                                            | 15/01/2020 | EN ISO 11607-1:2017          |
| 16 | ISO 11607-1: 2019 Packaging for terminally sterilized medical devices                                                                                                                                                                            | 02.2019    | ISO 11607-1: 2016/Amd 1:2014 |
| 17 | EN ISO 11607-2:2020 Packaging for terminally sterilized medical devices - Part 2: Validation requirements (ISO 11607-2: 2019)                                                                                                                    | 15.01.2020 | EN ISO 11607-2:2017          |
| 18 | ISO 11607-2: 2019 Packaging for terminally sterilized medical devices                                                                                                                                                                            | 02.2019    | ISO 11607-2: 2016/Amd 1:2014 |
| 19 | EN ISO 11737-1: 2018 Sterilization of medical devices - Microbiological methods - Part 1: Determination of a population of microorganisms on products (ISO 11737-1: 2018)                                                                        | 31.1.2018  | EN ISO 11737-1: 2006         |
| 20 | ISO 11737-2:2019 Sterilization of health care products — Microbiological methods — Part 2: Tests of sterility performed in the definition, validation and maintenance of a sterilization process                                                 | 2019-12    | ISO 11737-2:2009             |
| 21 | EN ISO 14937: 2009 Sterilization of health care products - General requirements for characterization of a sterilizing agent and the development, validation and routine control of a sterilization process for medical devices (ISO 14937: 2009) | 7.7.2010   | EN ISO 14937: 2000           |
| 22 | EN ISO 14971: 2019 Medical devices - Application of risk management to medical devices                                                                                                                                                           | 18.12.2019 | ISO 14971: 2012              |
| 23 | 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                                               | 20.11.2019 | EN ISO 11135:2014            |
| 24 | IEC 62366-1: 2015/Amd 1:2020 Medical devices – Part 1: Application of usability engineering to medical devices                                                                                                                                   | 17/06/2020 | IEC 62366-1: 2007/Amd 1:2014 |
| 25 | EN ISO 13485:2016 Medical devices - Quality management systems - Requirements for regulatory purposes (ISO 13485:2016)                                                                                                                           | 02/03/2016 | EN ISO 13485: 2012           |
| 26 | EN ISO 13485:2016/AC:2018                                                                                                                                                                                                                        | 28.03.2018 | ISO 13485:2016               |
| 27 | MEDDEV 2.7/1 Revision 4, Clinical evaluation, a Guide for manufacturers and notified bodies, under directives 93/42/EEC and 90/385/EEC                                                                                                           | 01.7.2016  | MEDDEV 2.7/1 Revision 3      |
| 28 | EN ISO 14644-1-2015 Cleanrooms and associated controlled environments — Part 1: Classification of air cleanliness by particle concentration                                                                                                      | 23/12/2015 | EN ISO 14644-1-1999          |
| 29 | EN 1644 -1:1997 Test methods for nonwoven compresses for medical use - Part 1: Nonwovens used in the manufacture of compresses                                                                                                                   | 19/02/1997 | First publication            |