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## EC Certificate

## **Production Quality Assurance System**

Directive 93/42/EEC on Medical Devices (MDD), Annex V (Devices in Class IIa, IIb or III)

No. G2 16 07 78455 013

Category(ies):

Manufacturer: Zhanjiang Star Enterprise Co., Ltd.

No. 1, West Jinhua Road

Mazhang District 524094 Zhanjiang

PEOPLE'S REPUBLIC OF CHINA



Corp. GmbH (Europe)

Eiffestraße 80 20537 Hamburg GERMANY

Product Foley Catheter (latex), Nelaton Catheter (latex),

Endotracheal Tubes, Anesthesia Kits,

**Suction Catheter for Single Use** 

in the Respiratory Tract,

**Disposable Urethral Catheterization Set,** 

Urethral Catheters (PVC), Stomach Tube (PVC),

Reinforced Endotracheal Tube

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex V. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class IIb and III devices an additional Annex III certificate is mandatory. See also notes overleaf.

Report No.: SH16090EXT01

**Valid from:** 2016-09-17 **Valid until:** 2021-09-16

Date. 2016-08-23

Stefan Preiß

04052768093615

TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

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# EC Certificate Production Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex V (Devices in Class IIa, IIb or III)

No. G2 16 07 78455 013

Facility(ies):

Zhanjiang Star Enterprise Co., Ltd.

No. 1, West Jinhua Road, Mazhang District, 524094

Zhanjiang, PEOPLE'S REPUBLIC OF CHINA

Zhanjiang Star Enterprise Co., Ltd.

No. 49 Jinchuan Road, 524094 Zhanjiang,

PEOPLE'S REPUBLIC OF CHINA



## EC Certificate

### **Production Quality Assurance System**

Directive 93/42/EEC on Medical Devices (MDD), Annex V (Devices in class I in sterile conditions, sterilised systems or procedure packs)

No. G2S 15 06 78455 011

Manufacturer: Zhanjiang Star Enterprise Co., Ltd.

> No. 1, West Jinhua Road Mazhang District

524094 Zhanjiang

PEOPLE'S REPUBLIC OF CHINA

**EC-Representative:** Shanghai International Holding

Corp. GmbH (Europe)

Eiffestraße 80 20537 Hamburg **GERMANY** 

**Product** Male External Catheter

Category(ies): Male External Catheter(Silicone)

> /Silicone Condom Catheter, **Endotracheal Tube Holder**, Mouth Gag (mouth opener),

**Drainage Bag** 

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture in accordance with MDD Annex V. This quality assurance system covers those aspects of manufacture concerned with securing and maintaining sterile conditions of the respective devices / device categories and conforms to the requirements of this Directive. It is subject to periodical surveillance. See also notes overleaf.

Report No.: SH15090EXT01

Valid from: 2015-09-16

Valid until: 2020-09-15

2015-07-14 Date,

Hans-Heiner Junker

TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

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#### **EC** Certificate

#### **Production Quality Assurance System**

Directive 93/42/EEC on Medical Devices (MDD), Annex V (Devices in class I in sterile conditions, sterilised systems or procedure packs)

No. G2S 15 06 78455 011

Facility(ies):

Zhanjiang Star Enterprise Co., Ltd.

No. 1, West Jinhua Road, Mazhang District, 524094

Zhanjiang, PEOPLE'S REPUBLIC OF CHINA

Zhanjiang Star Enterprise Co., Ltd.

No. 49 Jinchuan Road, 524094 Zhanjiang,

PEOPLE'S REPUBLIC OF CHINA

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**Product Service** 

# EC Certificate

### **Full Quality Assurance System**

Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4) (Devices in Class IIa, IIb or III)

No. G1 14 06 78455 010

Manufacturer:

Zhanjiang Star Enterprise Co., Ltd.

No. 1, West Jinhua Road **Mazhang District** 524094 Zhanjiang

PEOPLE'S REPUBLIC OF CHINA

**EC-Representative:** 

**Shanghai International Holding** 

Corp. GmbH (Europe)

Eiffestraße 80 20537 Hamburg **GERMANY** 

**Product** Category(ies):

Urethral Catheters (silicone), **Tracheostomy Tubes for Single Use** 

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

Report No.:

SH1409014

Valid from:

2014-09-16 2019-09-15

Valid until:

Hans-Heiner Junker



TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

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Date, 2014-09-17

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**EC** Certificate

**Full Quality Assurance System** 

Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4) (Devices in Class IIa, IIb or III)

No. G1 14 06 78455 010

Facility(ies):

Zhanjiang Star Enterprise Co., Ltd.

No. 1, West Jinhua Road, Mazhang District, 524094

Zhanjiang, PEOPLE'S REPUBLIC OF CHINA

Zhanjiang Star Enterprise Co., Ltd.

No. 49 Jinchuan Road, 524094 Zhanjiang,

PEOPLE'S REPUBLIC OF CHINA