

### Zhejiang Gongdong Medical Technology Co., Ltd

ADD: No.10, Belyuan Ave., Huangyan, Taizhou, Zhejiang, China, 318020

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### **AUTHORIZATION LETTER**

TO WHOM IT MAY CONCERN

MANUFACTURER

Zhejiang Gongdong Medical Technology Co., Ltd No.10, Beiyuan Ave., Huangyan, Taizhou, Zhejiang, China, 318020 DO HEREBY AUTHORIZE

"Echipamed Plus" SRL

Valea Trandafirilor 24B, of.80, MD-2001, Chisinau, Republic of Moldova
As our distributor and representative in Republic of Moldova for the medical disposable plastic ware products produced by Zhejiang Gongdong Medical Technology Co., Ltd.

This authorization letter valid from 18th, May, 2018 to 17th, May, 2021

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Zhejiang Gongdong Medical Technology Co., Ltd

Date 2018-05-18

Jim Qiu

Sales Manage





CRT2 / 10.13

DAKKS



# CERTIFICATE

No. Q1N 16 01 42464 026

**Holder of Certificate: Zhejiang Gongdong Medical** 

Technology Co., Ltd.

No.10 Beiyuan Ave., Huangyan 318020 Taizhou, Zheijang

PEOPLE'S REPUBLIC OF CHINA

Certification Mark:





Scope of Certificate:

Design and Development, Production and Distribution of Plastic Centrifuge Tubes,

Plastic Pipette Tips, Plastic Culture Dishes,

Plastic Forceps,

Plastic Test Tubes, Plastic Sample Cups,

Plastic First Aid Cases and Disposable Vacuum Blood Tubes,

Disposable Vacuum Blood Collection Systems,

Disposable Vaginal Speculum, Disposable Sterile Swabs,

Transportation Swabs with Medium. Micro Blood Collection Tubes. Capillary Blood Collection Tubes, Vacuum Urine Collection Sets, **Disposable Umbilical Cord Scissors** 

Disposable Specimen Container, Needle Holder.

Disposable Non Vacuum Blood tubes,

Disposable Anoscope, Disposable Loop Stick,

Sterile Vaginal Applicator

The Certification Body of TÜV SÜD Product Service GmbH certifies that the company mentioned above has established and is maintaining a quality management system, which meets the requirements of the listed standard(s). See also notes overleaf.

Report No.:

SH1611117

Valid from:

2016-02-29

Valid until:

2019-02-28

Date, 2016-02-26

Stefan Preiß

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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 16 01 42464 029

Manufacturer:

**Zhejiang Gongdong Medical** 

Technology Co., Ltd.

No.10 Beiyuan Ave., Huangyan 318020 Taizhou, Zhejiang

PEOPLE'S REPUBLIC OF CHINA



**EC-Representative:** 

Shanghai International Holding

Corp. GmbH (Europe)

Eiffestraße 80 20537 Hamburg **GERMANY** 

**Product** 

Category(ies):

Disposable Plastic Forceps, Disposable Vaginal Speculum,

Disposable Sterile Swabs,

Transportation Swabs with Medium,

Disposable Anoscope, Sterile Vaginal Applicator

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.:

SH1611117

Valid from:

2016-02-29

Valid until:

2020-09-28

Date, 2016-02-26

Stefan Preiß



TÜV SÜD Product Service GmbH is Notified Body with identification no Page 1 of 2

TÜV SÜD Product Service GmbH - Zertifizierstelle - Ridlerstraße 65 - 80339 Münshen YGermany

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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 18 03 42464 031

Manufacturer:

**Zhejiang Gongdong Medical** 

Technology Co., Ltd.

No.10 Beiyuan Ave., Huangyan 318020 Taizhou, Zhejiang

PEOPLE'S REPUBLIC OF CHINA



EC-Representative:

Shanghai International Holding

Corp. GmbH (Europe)

Eiffestraße 80 20537 Hamburg **GERMANY** 

**Product** Category(ies):

Disposable Vacuum Blood Collection System,

Disposable Umbilical Cord Scissors

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.:

SH18111EXT01

Valid from:

2018-06-09

Valid until:

2023-06-08

Date, 2018-04-05

Stefan Preiß

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

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