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Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
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ZLG-BS-244.10.08

TÜV SÜD Product Service

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 042464 2434

Manufacturer

Zhejiang Skg Medical Technology Co.,Ltd
No.39 Anye Road., Huangyan
318020 Taizhou, Zhejiang
PEOPLE'S REPUBLIC OF CHINA

Product Category(ies):

Disposable Vacuum Blood Collection System,
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.: SH19111EX901

Valid from: 2019-11-04
Valid until: 2024-05-26

Date, 2019-11-04

Christoph Dicks
Head of Certification/Notified Body

Digitally signed by Guțan Ghenadie
Date: 2021.10.01 13:35:06 EEST
Reason: MoldSign Signature
Location: Moldova

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

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Facility(ies):

Zhejiang Skg Medical Technology Co.,Ltd
No.39 Anye Road, Huangyan, 318020 Taizhou, Zhejiang, PEOPLE'S REPUBLIC OF CHINA

Zhejiang Skg Medical Technology Co.,Ltd
No.39 Anye Road, Huangyan, 318020 Taizhou, Zhejiang, PEOPLE'S REPUBLIC OF CHINA

ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

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