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ZLG-BS-244.10.08

Product Service

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 072857 0013 Rev. 00

Digitally signed by Moraru Grigore
Date: 2025.08.21 10:30:18 EEST
Reason: MoldSign Signature
Location: Moldova
MOLDOVA EUROPEANĂ

Manufacturer:

Changzhou Jiafeng Medical Equipment Co., Ltd.
Ninghe Village, Zhenglu Town
Tianning District
213115 Changzhou City, Jiangsu Province
PEOPLE'S REPUBLIC OF CHINA

Product Category(ies):

Infusion Sets for Single Use (with Needle), Sterile Hypodermic Syringes for Single Use (with Needle), Disposable Sterile Needles

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.: SH19603EXT01

Valid from: 2019-10-11
Valid until: 2024-05-26

Date, 2019-10-11

Stefan Preiß
Head of Certification/Notified Body

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

TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany

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