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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 103129 0002 Rev. 00

Manufacturer:

Shandong Chengwu Medical Products Factory
Southern end of Quancheng Road
Chengwu County
274200 Heze City, Shandong Province
PEOPLE'S REPUBLIC OF CHINA

Product Category(ies):

Disposable sterile venous blood specimen collection needle, Disposable infusion set.

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.: BJ1776601_BJ19766011

Valid from: 2019-12-06
Valid until: 2024-02-22

Date, 2019-12-06

Christoph Dicks
Head of Certification/Notified Body

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

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