



Benannt durch/Designated by
 Zentralstelle der Länder
 für Gesundheitsschutz
 bei Arzneimitteln und
 Medizinprodukten
 www.zlg.de
 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 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
 ZERTIFIKAT ◆ CERTIFICATE ◆ 認 證 證 書 ◆ CERTIFICADO ◆ CERTIFICADO ◆ CERTIFICAT



Benannt durch/Designated by
 Zentralstelle der Länder
 für Gesundheitsschutz
 bei Arzneimitteln und
 Medizinprodukten
 www.zlg.de
 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 103129 0002 Rev. 00

Facility(ies):

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

TÜV SÜD
 ZERTIFIKAT ◆ CERTIFICATE ◆ 認證書 ◆ CERTIFICADO ◆ CERTIFICAT

A4 / 07.17