



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 081232 0010 Rev. 01

Manufacturer:

**AnHui Hongyu Wuzhou
Medical Manufacturer Co.,Ltd.**

No.2 Guanyin Road
Economic Development Zone
Taihu County
246400 Anqing, Anhui
PEOPLE'S REPUBLIC OF CHINA

EC-Representative:

Shanghai International Holding Corp. GmbH (Europe)
Eiffestraße 80, 20537 Hamburg, GERMANY

**Product
Category(ies):**

**Sterile Hypodermic Syringes for Single Use, Sterile
Hypodermic Needles for Single Use, Sterile Insulin Syringes
for Single Use, Insulin Pen-injectors for Medical Use,
Transfusion Sets for Single Use, Infusion Sets for Single
Use(Gravity Feed), Burette-type Infusion Sets for Single Use,
Intravenous Needles for Single Use, Blood Collection Needles
for Single Use, Sterile Dental Injection Needles for Single
Use, Disposable Precision Flow Regulator, Insulin Pen
Needles for Single Use, Blood Collection Sets for Single Use,
Sterile Safety Hypodermic Needles for Single Use, Safety
Blood Collection Sets for Single Use, Safety Blood Collection
Needles for Single Use**

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.:

SH1873508

Valid from:

2018-08-01

Valid until:

2022-08-19

Date,

2018-08-01

Stefan Preiß

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ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ CERTIFICADO ◆ CERTIFICAT



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