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

Full Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in Class IIa, IIb or III)

No. G1 001460 0002 Rev. 01

Manufacturer:

Shenzhen Maiwei Biotech Co., Ltd.
5/F, Office Building, 2/F, Building 1
2-10 Jinlong Blvd. South
Pingshan District
518118 Shenzhen
PEOPLE'S REPUBLIC OF CHINA

Product Category(ies): **High Pressure Syringe and Pressure Connecting Tube**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

Report No.: GZ1931401

Valid from: 2020-02-13
Valid until: 2022-11-16

Date, 2020-02-13

Christoph Dicks
Head of Certification/Notified Body

Podpisany elektronicznie przez Katarzyna Paulina Cieśla
22.06.2023 12:49:02 +02'00'

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

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

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Facility(ies):

Shenzhen Maiwei Biotech Co., Ltd.
5/F, Office Building, 2/F, Building 1, 2-10 Jinlong Blvd. South,
Pingshan District, 518118 Shenzhen, PEOPLE'S REPUBLIC OF CHINA

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Podpisany elektronicznie przez
Katarzyna Paulina Cieśla
22.06.2023
12:49:21 +02'00'

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TÜV SÜD