



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 17 08 01460 002

Manufacturer: **Shenzhen Maiwei Biotech Co., Ltd.**

2/F, Building 1, 2-10 Jinlong Blvd. South
Pingshan District
518118 Shenzhen
PEOPLE'S REPUBLIC OF CHINA



EC-Representative: **Shanghai International Holding Corp. GmbH (Europe)**

Eiffestraße 80
20537 Hamburg
GERMANY

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.

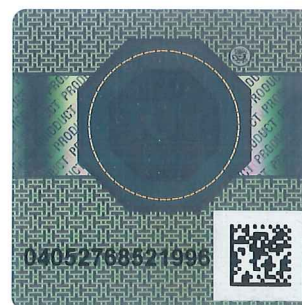
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Date, 2017-11-17

S. Preiß

Stefan Preiß



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

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Facility(ies):

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