



Product Service

# CERTIFICATE

No. Q5 17 08 01460 001

**Holder of Certificate:** Shenzhen Maiwei Biotech Co., Ltd.

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

**Facility(ies):**

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



**Certification Mark:**



**Scope of Certificate:** Design and Development, Production and Distribution of High Pressure Syringe and Pressure Connecting Tube

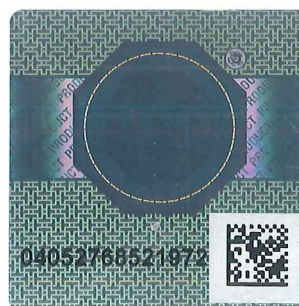
**Applied Standard(s):**

EN ISO 13485:2016  
Medical devices - Quality management systems -  
Requirements for regulatory purposes  
(ISO 13485:2016)  
DIN EN ISO 13485:2016

The Certification Body of TÜV SÜD Product Service GmbH certifies that the company mentioned above has established and is maintaining a quality management system, which meets the requirements of the listed standard(s). See also notes overleaf.

**Report No.:** GZ1731401

**Valid from:** 2017-11-17  
**Valid until:** 2020-11-16



**Date,** 2017-11-17

*S. Preiß*  
Stefan Preiß

