

## 0

Rev. 00

No. Q6 050079 0013

Holder of

Certificate:

Facility(ies):

Co., Ltd. Shenzhen Ant Medical Devices

18 Jinhui Ave., Pingshan New Dist 518122 Shenzhen PEOPLE'S REPUBLIC OF CHINA Pingshan New District

Shenzhen Ant Medical Devices Co., Ltd.

18 Jinhui Ave., Pingshan New District, 518122 Shenzhen
PEOPLE'S REPUBLIC OF CHINA

Certification Wark



Scope of Certificate:

and Injection Tubing System. Pressure Syringe, Pressure Connecting Tube, Production, Sales and Distribution of High

Applied Standard(s):

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

DIN EN ISO 13485:2016

above has established and is maintaining a quality management system (excluding subclause 7.3), which meets the requirements of the listed standard(s). See also notes overleaf. The Certification Body of TÜV SÜD Product Service GmbH certifies that the company mentioned

Report No.:

BJ18899041

Valid until: Valid from:

2018-09-01 2021-08-31

2018-08-28

Date,

Stefan Preiß



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СЕРТИФИКАТ ◆

ZLG-BS-244.10.08

## PATICATE

(Devices in Class IIa, IIb or III) Directive 93/42/EEC on Medical Devices (MDD), Annex V Production Quality Assurance System

No. G2 050079 0012 Rev. 01

Manufacturer:

Shenzhen Ant Medical Devices

Co., Ltd.

18 Jinhui Ave., Pingshan New District

518122 Shenzhen
PEOPLE'S REPUBLIC OF CHINA

Category(ies): Product

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

EC-Representative:

Injection Tubing System. High Pressure Syringe, Pressure Connecting Tube,

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

Report No.:

認證證書

BJ1889907

Valid until: Valid from:

2023-09-17 2018-09-18

Date

CERTIFICATE

2018-08-07

Stefan Preiß





ZERTIFIKAT ◆

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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Production Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex V
(Devices in Class IIa, IIb or III)

No. G2 050079 0012 Rev. 01

Facility(ies):

Shenzhen Ant Medical Devices Co., Ltd.
18 Jinhui Ave., Pingshan New District, 518122 Shenzhen, PEOPLE'S REPUBLIC OF CHINA

