

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 15 08 73650 010

Manufacturer:

Shenzhen Probe

Science & Technology Co., Ltd.

4th Floor

6th Floor and Zone A of 9th Floor Xintian Road, Fuyong Street

Block C, No. 71-3 Bao'an District

518103 Shenzhen, Guangdong PEOPLE'S REPUBLIC OF CHINA

EC-Representative:

Wellkang Ltd

Suite B, 29 Harley Street

LONDON W1G 9QR

UNITED KINGDOM

Product

Category(ies):

Anesthesia Machine, Ventilator, Air Compressor, Respiratory Humidifier, Vaporizer, Emergency and

Transport Ventilator, Full Digital Ultrasonic

Diagnostic Instrument

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.:

7484007379

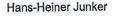
Valid from:

2015-10-20

Valid until:

2020-10-12

Date, 2015-11-19





TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123 Page 1 of 2



Product Service

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Facility(ies):

Shenzhen Probe Science & Technology Co., Ltd. 4th Floor, 6th Floor and Zone A of 9th Floor, Xintian Road, Fuyong Street, Block C, No. 71-3, Bao'an District, 518103 Shenzhen, Guangdong, PEOPLE'S REPUBLIC OF CHINA

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