ZERTIFIKAT ◆ CERTIFICATE ◆ 認証証書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

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 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

Hans-Heiner Junker

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

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Facility(ies):

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Fuyong Street, Block C, No. 71-3, Bao`an District, 518103
Shenzhen,Guangdong, PEOPLE'S REPUBLIC OF CHINA