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Zentralstelle der Länder
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bei Arzneimitteln und
Medizinprodukten
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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 057571 0003 Rev. 00

Manufacturer:

**Beijing Choice Electronic
Technology Co., Ltd.**

2nd Floor
3rd Floor and Room 410-412 4th Floor
No. 2 Building, No. 9 Shuangyuan Road
Shijingshan District
100041 Beijing
PEOPLE'S REPUBLIC OF CHINA

Product Category(ies): **Pulse Oximeter, Vital Sign Monitor, Pulse Oximeter Sensor, Handheld ECG Monitor, Fetal Doppler, Fingertip Pulse Oximeter with Forehead Thermometer, Wireless Thermometer, Blood Pressure Monitor, Handheld Multi-parameter Patient Monitor.**

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.: BJ19901031

Valid from: 2020-03-27
Valid until: 2024-05-21

Date, 2020-03-27

Christoph Dicks
Head of Certification/Notified Body

TÜV SÜD
 ZERTIFIKAT ◆ CERTIFICATE ◆ 認證書 ◆ CERTIFICADO ◆ CERTIFICAT

