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Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
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TÜV SÜD 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 078179 0032 Rev. 01

Manufacturer:
Beijing Choice Electronic Technology Co., Ltd.
Room 4104, No. A12 Yuquan Road
Haidian District
100143 Beijing
PEOPLE'S REPUBLIC OF CHINA

EC-Representative:
Shanghai International Holding Corp. GmbH (Europe)
Eiffestraße 80, 20537 Hamburg, GERMANY

Product Category(ies):
Portable Patient Monitor, Pulse Oximeter, Vital Sign Monitor, Pulse Oximeter Sensor, Handheld ECG Monitor, Fetal Doppler, Fingertip Pulse Oximeter with Forehead Thermometer, Compressor Nebulizer, 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.

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Valid from: 2019-05-22
Valid until: 2024-05-21

Date, 2019-05-08

Stefan Preiß

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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No. G1 078179 0032 Rev. 01

Facility(ies):

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