Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
ZLG-BS-244.10.08

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 092582 0009 Rev. 00

Manufacturer: **Xuzhou Yongkang Electronic Science Technology Co., Ltd.**
4F Building C8, 40 Jingshan Road
Economic and Technological Development Zone
221000 Xuzhou
PEOPLE'S REPUBLIC OF CHINA

Facility(ies): Xuzhou Yongkang Electronic Science Technology Co., Ltd.
4F Building C8, 40 Jingshan Road, Economic and Technological Development Zone, 221000 Xuzhou, PEOPLE'S REPUBLIC OF CHINA

Product Category(ies): **Vital Signs Monitor,**
Handheld Pulse Oximeter,
Fingertip Pulse Oximeter,
Multiparameter 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.: SH1966201

Valid from: 2019-11-26
Valid until: 2024-05-26

Date, 2019-11-26

Christoph Dicks
Head of Certification/Notified Body

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

TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany

ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT
TÜV SÜD
A4 / 07.17