ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

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 17 11 63196 017

Manufacturer:

Shenzhen Hexin ZONDAN Medical Equipment Co., Ltd.
Floor 14, Block D
Dianlian Technology Building
the Crossing between South Circle Road and South Fuli Road
Guangming District
518106 Shenzhen
PEOPLE'S REPUBLIC OF CHINA

EC-Representative:

Shanghai International Holding Corp. GmbH (Europe)
Eiffestraße 80
20537 Hamburg
GERMANY

Product Category(ies):

Pulse Oximeter,
Patient Monitor,
Fetal Dopplers,
Fetal 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.: SH1722301

Valid from: 2018-04-11
Valid until: 2023-04-10

Date, 2018-04-11

Stefan Preiß

04052768706911

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

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No. G1 17 11 63196 017

Facility(ies):

Shenzhen Hexin ZONDAN Medical Equipment Co., Ltd.
Floor 14, Block D, Dianlian Technology Building,
the Crossing between South Circle Road and
South Fuli Road, Guangming District, 518106
Shenzhen, PEOPLE'S REPUBLIC OF CHINA

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