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Zentralstelle der Länder  
für Gesundheitsschutz  
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 044751 0167 Rev. 02**

**Manufacturer: Shenzhen Mindray Bio-Medical Electronics Co., Ltd.**

Mindray Building  
Keji 12th Road South  
High-Tech Industrial Park  
Nanshan  
518057 Shenzhen  
PEOPLE'S REPUBLIC OF CHINA

**Product Category(ies):** Patient Monitoring Devices,  
Vital Signs Monitor,  
Center Monitoring System,  
Telemetry Monitoring System,  
Ambulatory Blood Pressure Monitor,  
Pulse Oximeter, Temperature Probe,  
SPO2 Sensors, Electrocardiograph,  
Ventilator, Anesthetic Vaporizer,  
Air compressor,  
Ultrasonic Diagnostic Equipment,  
Ultrasonic Transducer,  
Digital Radiography System,  
Radiography System

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:** 2019-11-13  
**Valid until:** 2024-05-26

**Date,** 2019-11-13

Christoph Dicks  
Head of Certification/Notified Body

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

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 ZERTIFIKAT ◆ CERTIFICATE ◆ 認證書 ◆ CERTIFICADO ◆ CERTIFICAT

