



Benannt durch/Designated by  
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 070744 0017 Rev. 00**

**Manufacturer:**

**Nanjing Mindray Bio-Medical  
Electronics Co., Ltd.**

666# Middle Zhengfang Road

Jiangning

211111 Nanjing, Jiangsu

PEOPLE'S REPUBLIC OF CHINA

**Product Category(ies): Medical Supply Units**

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.:**

SH2055901

**Valid from:**

2020-03-05

**Valid until:**

2024-05-26

**Date,**

2020-03-05

Christoph Dicks

Head of Certification/Notified Body