



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 10 49957 033

**Manufacturer:**

**Guangdong Biolight  
Meditech Co., Ltd.**

No.2 Innovation First Road  
Technical Innovation Coast  
Hi-tech Zone, Zhuhai  
519085 Zhuhai, Guangdong  
PEOPLE'S REPUBLIC OF CHINA



**EC-Representative:**

**Shanghai International Holding  
Corp. GmbH (Europe)**

Eiffestraße 80  
20537 Hamburg  
GERMANY

**Product  
Category(ies):**

**Patient Monitor, Fetal Monitor,  
Central Monitoring System Software,  
Pulse Oximeter, Electrocardiograph,  
Electronic Thermometer,  
Electronic Sphygmomanometer,  
Ultrasonic Doppler Fetal Heartbeat Detector,  
Syringe Pump used for intravenous injection  
administration,  
Infusion Pump used for intravenous infusion  
administration,  
SpO2 Sensors, Temperature Probes**

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.:**

SH1725816

**Valid from:**

2018-04-16

**Valid until:**

2020-03-19

**Date,** 2018-04-16

Stefan Preiß



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

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**Facility(ies):**

Guangdong Biolight Meditech Co., Ltd.  
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Coast, Hi-tech Zone, Zhuhai, 519085 Zhuhai,  
Guangdong, PEOPLE'S REPUBLIC OF CHINA