



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 14 09 78179 017

**Manufacturer:** **Beijing Choice Electronic  
Technology Co., Ltd.**  
Room 4104, No. A12 Yuquan Road  
Haidian District  
100143 Beijing  
PEOPLE'S REPUBLIC OF CHINA

**EC-Representative:** **Shanghai International Holding  
Corp. GmbH (Europe)**  
Eiffestraße 80  
20537 Hamburg  
GERMANY

**Product  
Category(ies):** **Portable Patient Monitor, Pulse Oximeter,  
Vital Sign Monitor, Pulse Oximeter Sensor,  
Handheld ECG Monitor, Fetal Doppler,  
Compressor Nebulizer,  
Fingertip Pulse Oximeter with Forehead Thermometer.**

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.:** BJ1490104-3

**Valid from:** 2014-11-18

**Valid until:** 2019-05-21



**Date,** 2014-11-19

Hans-Heiner Junker

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

Page 1 of 2





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**Facility(ies):**

Beijing Choice Electronic Technology Co., Ltd.  
Room 4104, No. A12 Yuquan Road, Haidian District, 100143  
Beijing, PEOPLE'S REPUBLIC OF CHINA

Beijing Choice Electronic Technology Co.,Ltd.  
Floor 4, Jingyang Building, No.15, Xijing Rd., Shijingshan District,  
100041 Beijing, PEOPLE'S REPUBLIC OF CHINA

Beijing Choice Electronic Technology Co., Ltd.  
No.9 Shuangyuan Rd., Badachu Hi-tech Zone, Shijingshan  
District, 100041 Beijing, PEOPLE'S REPUBLIC OF CHINA