



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 13 04 78179 007

Manufacturer:

**Beijing Choice Electronic
Technology Co., Ltd.**

Room 320, West Building 4, No.83 Fuxing Road
100039 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.**

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.:

BJ1390104-1

Valid from:

2013-10-29

Valid until:

2014-05-21



Date, 2013-10-31

Hans-Heiner Junker

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

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No. G1 13 04 78179 007**Facility(ies):**

Beijing Choice Electronic Technology Co., Ltd.
Room 320, West Building 4, No.83 Fuxing Road, 100039 Beijing,
PEOPLE'S REPUBLIC OF CHINA

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