



Product Service

EC - CERTIFICATE

Full Quality Assurance System

(Annex II, section 3 of the Directive 93/42/EEC on Medical Devices)

No. G1 09 04 50247 012

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

Bailangyuan Building B
Rm. 1127-1128, Fuxing Road, A36
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**

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 products / product categories according to Annex II section 3 of the Directive 93/42/EEC on Medical Devices. This quality assurance system conforms to the provisions of this Directive and is subject to periodical surveillance. For marketing of class III products an additional Annex II.4 certificate is mandatory. See also notes overleaf.

Report No.: BJ890104

Valid until: 2014-05-21

Date, 2009-05-22

Hans-Heiner Junker



TÜV SÜD Product Service GmbH is Notified Body according to Council Directive 93/42/EEC concerning medical devices with identification no. 0123.

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Facility(ies):

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100039 Beijing, PEOPLE'S REPUBLIC OF CHINA

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