

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



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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Date.

2009-05-22

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Facility(ies):

Beijing Choice Electronic Technology Co., Ltd. Bailangyuan Building B, Rm. 1127-1128, Fuxing Road, A36, 100039 Beijing, PEOPLE'S REPUBLIC OF CHINA

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