



Product Service

EC - CERTIFICATE

Full Quality Assurance System

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

No. G1 10 05 49076 003

Manufacturer: **Shenzhen Creative Industry Co., Ltd.**

2/F, Block 3
Nanyou Tian'an Industry Town
518054 Shenzhen, GD
PEOPLE'S REPUBLIC OF CHINA

EC-Representative: **Shanghai International Trading Corp. GmbH (Hamburg)**

Eiffestrasse 80
20537 Hamburg
GERMANY

Product Category(ies): **Patient Monitor, Vital Signs Monitor, Fetal Doppler, Fingertip Oximeter, Handheld Pulse Oximeter, Wrist Oximeter, Easy 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.: SZ090430101

Valid until: 2015-10-12

Date, 2010-10-13

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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