ZERTIFIKAT ◆ CERTIFICATE ◆ 認証証書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

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 10 12 61623 006

Manufacturer:
Meditech Equipment Co., Ltd.
1003 Jinhua Plaza, 100# Nanjing Road
266071 Qingdao, Shandong Province
PEOPLE'S REPUBLIC OF CHINA

EC-Representative:
FARAG LTD
Klearchou 7 - Galatsi
11147 Athens
GREECE

Product Category(ies):
Pulse Oximeter, Electrocardiography, Visual Stethoscope, Fetal Doppler, Fetal Monitor, B-Ultrasound Diagnostic System

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.: SH1008803

Valid from: 2011-06-07
Valid until: 2012-04-29

Date, 2011-06-10

Hans-Heiner Junker

TÜV SÜD 479094

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

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Facility(ies):

Meditech Equipment Co., Ltd.
1003 Jinhua Plaza, 100# Nanjing Road, 266071
Qingdao, Shandong Province, PEOPLE'S REPUBLIC OF CHINA

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