



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 18 03 66456 020

Category(ies):

Manufacturer: **Unimed Medical Supplies, Inc.**

Bld#8, Nangang 3rd Industrial Park

Tangtou, Shiyan 518108 Shenzhen

PEOPLE'S REPUBLIC OF CHINA

Facility(ies): Unimed Medical Supplies, Inc.

Bld#8, Nangang 3rd Industrial Park, Tangtou, Shiyan, 518108

Shenzhen, PEOPLE'S REPUBLIC OF CHINA

Product Temperature Probe for Medical Use, SpO2 Sensor,

Ultrasonic Fetal Heart Detector, Oximeter,

Electrocardiograph, Disposable Blood Pressure Transducers, Oxygen Sensor for Medical Use, Fetal

transducer

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.: GZ18134EXT01

Valid from: 2018-06-09

Valid until: 2023-06-08

Date, 2018-06-06 Stefan Preiß



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

Page 1 of 1