



Product Service

EC Certificate

Production Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex V
(Devices in Class IIa, IIb or III)

No. G2 16 10 97295 002

Manufacturer: **WAT Medical Technology Inc.**

Room 703-711, No.2 North Taoyuan Road
315600 Ningbo, Zhejiang Province
PEOPLE'S REPUBLIC OF CHINA



EC-Representative: **KINGSMEAD SERVICE LIMITED**

145-157 St John Street
London
EC1V 4PY
UNITED KINGDOM

Product Category(ies): **TENS-device**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex V. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class IIb and III devices an additional Annex III certificate is mandatory. See also notes overleaf.

Report No.: SH16112001

Valid from: 2017-06-16

Valid until: 2022-04-26



Date, 2017-06-16

Stefan Preiß

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

Page 1 of 2



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