

## EC Certificate

## Production Quality Assurance

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

No. G2 16 01 78453 008

**Wanufacturer:** 

Alich Medical (Shenzhen) Co., LTD.

3/F, No.4 Building

Niucheng Industrial Park, XIII

Nanshan District

518055 Shenzhen, Guangdong Province PEOPLE'S REPUBLIC OF CHINA

EC-Representative:

Renault-Petersen Limited

Couching House, Couching Street

Wallington

Oxfordshire OX49 5PX UNITED KINGDOM

Product

Category(ies):

Clinical Infrared Thermometer, Electronic
Thermometer, Nerve and Muscle Stimulator,
Ultrasound Physical Therapy Device, Continuos
Passive Motion Device, Radiofrequency Stimulators,
Cryos Therapy Device, Magneto Therapy Device,
Laser Therapy Device, Shock Wave Therapy 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 II excl. section 4. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance.

Report No.:

748442701

Valld from:

2016-10-10

Valid until:

2021-03-01

Date, 2016-10-10

Hans-Heiner Junker

382326

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

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