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Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
ZLG-BS-244.10.08

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 090072 0007 Rev. 01**

**Manufacturer:**

**SHENZHEN YINGCHI TECHNOLOGY CO., LTD**
Block B, 2/F
Juyou Entrepreneurship Center (Phoenix City Building)
15 Sci-tech North 1st Road
Songpingshan Community Xili
Nanshan District
518057 Shenzhen
PEOPLE'S REPUBLIC OF CHINA

**Product Category(ies): Pulsed Magnetic Stimulation Device**

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.:** GZ2019803

**Valid from:** 2021-03-06
**Valid until:** 2024-05-26

**Date,** 2021-03-02

Christoph Dicks
Head of Certification/Notified Body


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

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