EC Certificate

Directive 93/42/EEC Annex V
Production Quality Assurance
Medical Devices

Registration No.: DD 60129671 0001

Report No.: 17030157 007

Manufacturer:
EasyMed Instruments Co., Ltd.
3/F, 5/F- 6/F, Block A, Gupo Gongmao
Building, Fengxin Road, Fengxiang
Industrial District, Daliang,
528300 Shunde, Foshan, Guangdong
China

Products:
- Neuromuscular Stimulators
- Transcutaneous Electrical Nerve Stimulators
- TENS/EMS/ Micro current/ Interferential Stimulators
- Peripheral Nerve Stimulators
  Transcutaneous Vagus Nerve Stimulators (tVNS)

Replaces Approval, Registration No.: DD 60114968 0001

Expiry Date: 2023-04-09

The Notified Body hereby declares that the requirements of Annex V of the directive 93/42/EEC have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance, defined by Annex V, section 4 of the aforementioned directive. For placing on the market of class IIb and class III devices covered by this certificate an EC type-examination certificate according to Annex III is required.

Effective Date: 2018-08-15

Date: 2018-08-16

Notified Body

S. Liu

TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg

TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC concerning medical devices with the identification number 0197.