EC Certificate

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

Registration No.: DD 60109994 0001

Report No.: 15081806 002

Manufacturer:
VINNO Technology (Suzhou)
Co., Ltd.
5F Building A,4F Building C
No. 27 Xinfa Rd.
Suzhou Industrial Park
Suzhou
215123 Jiangsu
China

Products: Ultrasound Diagnostic Systems

Replaces Approval, Registration No.: DD 60101669 0001

Expiry Date: 2020-07-19

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: 2016-05-24

Date: 2016-05-24

Notified Body
TÜV Rheinland LGA Products GmbH
TÜVRheinland
Zertifizierstelle
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.

Digitally signed by Valiuh Tatiana
Date: 2020.05.28 15:00:25 EEST
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