EC Certificate

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

Registration No.: DD 60136869 0001

Report No.: 15064657 007

Manufacturer:
Wujiang City Shenlong Medical
Health Product Co., Ltd.
Gangzi Cun, Shenta, Lili Town
Wujiang District
Suzhou
215213 Jiangsu
China

Products:
- Disposable Acupuncture Needles
- Disposable Press Needles

Replaces the Approval, Registration No.: DD 60090111 0001

Digitally signed by Cobzari-Țurcan Rodica
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Expiry Date: 2024-01-17

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: 2019-04-19

Date: 2019-04-19

Notified Body

Fuxiu Sheng

TÜV Rheinland LGA Products GmbH Zertifizierungsstelle

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.