EC Certificate

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

Registration No.: DD 60138021 0001

Report No.: 15066755 007

Manufacturer:

Wujiang City Cloud &
Dragon Medical Device Co., Ltd.
Building 17, No. 148 Yucai Road
Beishe, Wujiang
Jiangsu Province, 215214
China

Products:

Disposable Acupuncture Needles

Replaces Approval, Registration No.: DD 60091366 0001

Expiry Date: 2024-02-18

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: 2019-05-09

Date: 2019-05-09

Notified Body

Fuxiu Sheng

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.