EC Certificate

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

Registration No.: DD 60141252 0001

Report No.: 16803369 008

Manufacturer:
Beijing Ruicheng Medical
Supplies Co., Ltd.
No. 558 Zhangzikou
Yangsong Town, Huairou District
101400 Beijing
China

Products: Medical Devices

(see attachment for products and additional site included)

Replaces Approval, Registration No.: DD 60094561 0001

Expiry Date: 2024-05-27

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: 2019-08-05

Date: 2019-08-05

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.

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TÜV Rheinland
LGA Products GmbH
Tillystraße 2, 90431 Nürnberg

Attachment to Certificate
Registration No.: DD 60141252 0001
Report No.: 16803369 008

Manufacturer:
Beijing Ruicheng Medical Supplies Co., Ltd.
No. 558 Zhangzikou
Yangsong Town, Huairou District
101400 Beijing
China

Products:

- Disposable Lancets

For the following medical devices the scope covers only the aspects of manufacture concerned with securing and maintaining sterile condition:

- Alcohol Prep Pads

Site included:

North Building, Fengxiang Development Zone, Zhangzikou Economic Cooperatives, Yangsong Town, Huairou District, 101400 Beijing, China

Beijing Ruicheng Medical Supplies Co.,Ltd. 1101160027307

Notified Body

TÜV Rheinland LGA Products GmbH Zertifizierungsstelle

Date: 2019-08-05

Fuxiu Sheng

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