

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 sites 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



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.

TÜV Rheinland
LGA Products GmbH
Tillystraße 2, 90431 Nürnberg

Doc. 1/1, Rev. 0

**Attachment to
Certificate**

Registration No.: DD 60141252 0001
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Products:

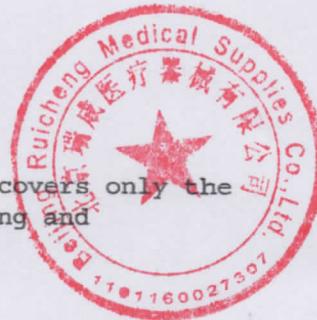
- Disposable Lancets

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



Date: 2019-08-05

Notified Body

Fuxiu Sheng

