

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

Registration No.: DD 60132306 0001

Report No.: 15096329 002

Manufacturer: Jiangsu Eyoung Medical Devices
Co., Ltd.
No. 1 Dongtang Road, Zhenglu Town
Tianning District
Changzhou
213115 Jiangsu
China

Products: Medical Devices
(see attachment for products included) ®

TÜVRheinland

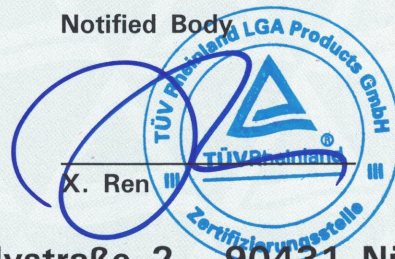
Expiry Date: 2023-09-10

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: 2018-09-11

Date: 2018-09-11

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

**Attachment to
Certificate**

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Manufacturer: Jiangsu Eyoung Medical Devices
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Products:

- Syringes for Single Use
- Disposable Hypodermic Needles

Aspects of manufacture concerned with securing and
maintaining sterile conditions:

- Infusion Spikes
- Oral Syringes (with or without Adaptors)

Date: 2018-09-11

