

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

Registration No.: DD 60150145 0001

Report No.: 15092196 007

Manufacturer: Yangzhou Beswin Medical
Equipment Co., Ltd.
NO.9, Zhongxing Rd.
Yuetang Town, Yizheng
Yangzhou
211413 Jiangsu
P.R. China

Products: Medical Devices

(see attachment for products included)

Expiry Date: 2024-05-26

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: 2020-06-23

Date: 2020-06-23

Notified Body

Herbert Zhong



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**

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Products:

- Endotracheal Tubes for single use
- Laryngeal Mask Airways for single use
- Breathing Circuits for single use
- Closed Suction Catheters for single use
- Breathing Filters for single use
- Anaesthesia Masks for single use
- Stomach Tubes for single use
- Disposable Endoscopic Retrieval Bags

Date: 2020-06-23

Notified Body

Herbert Zhong

