

TÜV Rheinland LGA Products GmbH • 51105 Köln

Ningbo Jiangbei Woson Medical Instrument Co., Ltd.  
No. 25, Lane 300, Jinshan Road,  
Jiangbei District, Ningbo  
315032 Zhejiang  
P.R. China

Contact

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Mail: [medical-products@de.tuv.com](mailto:medical-products@de.tuv.com)

Date March 25, 2024

**Application for: QMS**

Certificate No. : HD 2058015-1

Requirement : MDD 93/42/EEC Annex II excluding (4)

Dear Madame or Sir,

**Confirmation letter surveillance audit**

A surveillance audit of your quality management system was performed.

The audit team confirmed that your quality management system is applied effectively with respect to the above-mentioned requirements.

The recommendation of auditor is indicated in report no. 244565668-200

This letter confirms that the above-mentioned certificate will remain valid.

Best regards,



Jason Pan  
Certification body

TÜV Rheinland  
LGA Products GmbH

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51105 Köln  
Germany

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Board of Management

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Dr. Jörg Schlösser

Nuremberg HRB 26013  
VAT No.: DE 811835490

Chairman of the  
Supervisory Board

Dr.-Ing. Michael Fübi

# EC Certificate



**Full Quality Assurance System**  
**Directive 93/42/EEC on Medical Devices, Annex II excluding (4)**

Registration No.: HD 2058015-1

Manufacturer: Ningbo Jiangbei Woson Medical Instrument Co., Ltd.  
No. 25, Lane 300, Jinshan Road,  
Jiangbei District, Ningbo  
315032 Zhejiang  
P.R. China

Products: Steam Sterilizers

A large, light-colored watermark of the TÜVRheinland logo is centered on the page. It consists of a stylized triangle with a horizontal line through it, and the text 'TÜVRheinland' with a registered trademark symbol (®) to the right.

TÜVRheinland®

The Notified Body hereby declares that the requirements of Annex II, excluding section 4 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 II, section 5 of the aforementioned directive. For placing on the market of class III devices covered by this certificate an EC design-examination certificate according to Annex II section 4 is required.

Report No.: 244310228-200

Effective date: 2021-04-21

Expiry date: 2024-05-26

Issue date: 2021-04-21



A handwritten signature in blue ink that reads 'Jason Pan' is written over the seal and extends to the right.  
Jason Pan  
TÜV Rheinland LGA Products GmbH  
Tillystraße 2 · 90431 Nürnberg · Germany

TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC concerning medical devices with the identification number 0197.