



# EC Declaration of Conformity

*Manufacturer:*

**Ningbo Jiangbei Woson Medical Instrument Co., Ltd.**

No.25, Lane 300, Jinshan Road, Jiangbei District,  
Ningbo 315032, China  
Tel: 0086 574 83022668  
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*whose single Authorized Representative:*

**DTF TECHNOLOGY srl**

via Gressoney 9, 20137 Milano, Italy  
tel: +39.02.84893641  
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We, the manufacturer, herewith declare that the products

**Steam Sterilizer – TANDA STERI**

**UMDNS-Code: 13746; GMDN-Code: 38671**

meet the provisions of Directive 93/42/EEC and 2007/47/EEC which apply to them.

The medical device has been assigned to class IIb according to rule 15 Annex IX of the Directive 93/42/EEC. It bears the mark



The product concerned has been designed and manufactured under a quality management system according to Annex II (excluding Section 4) of Directive 93/42/EEC.

Compliance of the designated product with the Directive 93/42/EEC has been assessed and certified by the Notified Body

**TÜV Rheinland LGA Product GmbH -  
Tillystraße 2-90431 Nürnberg  
Identification Number: 0197**

**Certificate No.: HD 2058015-1**

**Issue date: 2021/04/21**

**Expiry date: 2024/05/26**

following the procedure relating to the EC Declaration of Conformity set out in Annex II (excluding Section 4) of Directive 93/42/EEC.

Applied harmonized standards, national standards or other normative documents:

EN 13060:2014, EN 61010-1:2010, EN 61010-2-040:2015, EN 61326-1:2018.

This Declaration of conformity is valid in connection with the release document for the respective batch of produced devices.

The above mentioned declaration of conformity is exclusively under the responsibility of

**Ningbo Jiangbei Woson Medical Instrument Co., Ltd.**

2021/04/28 – Ningbo, China

Place, date



Name: Xie Diyan

Position: Vice General Manager

Signature:

Legally binding signature, Function