

## EC Declaration of Conformity

Manufacturer:

whose single Authorized Representative:

Guilin Woodpecker Medical Instrument Co., Ltd.

MedNet EC-Rep GmbH • Borkstrasse 10 • 48163  
Muenster • Germany

We, the manufacturer, here with declare that  
the products **Ultrasonic Surgical System**, *UMDNS-Code: 18049*

Product name	Model
Piezo Bone Surgery	ULTRASURGERY、ULTRASURGERY LED
	US-II、US-II LED
	DS-II、DS-II LED
	Surgery-X、Surgery-X LED
	Surgic Touch、Surgic Touch LED
	Ai-surgery、Surgic smart

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

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

**CE 0197**

The product concerned has been manufactured under a quality management system according to Annex V and Annex VII 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 Products GmbH**  
**Tillystraße 2, 90431, Nürnberg, Germany**

Certificate No.: DD 60137494 0001

Issue date: 2019-7-16

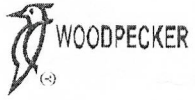
Expiry date: 2024-5-27

following the procedure relating to the EC Declaration of Conformity set out in Annex V and Annex VII of Directive 93/42/EEC.

This Declaration of Conformity covers all medical devices as specified in the product list belonging to this declaration and is only valid in connection with a batch specific Certificate of Compliance for all products concerned bearing the CE mark

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





桂林市啄木鸟医疗器械有限公司  
GUILIN WOODPECKER Medical Instrument Co.,LTD.

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Version:H

Company: Guilin Woodpecker Medical Instrument Co., Ltd.

Address: Information Industrial Park, Guilin National High-Tech Zone, Guilin, Guangxi,  
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陈燕 2021.12.20  
Preparation , Date

王进伟 2021.12.20  
Review , Date

于加波 2021.12.20  
Legally binding signature, Function

