

File No.: ZMN-YF-(E01)-01-001-02

Version: C

## 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, herewith declare that the products  
**Endo Motors UMDNS-Code: 35416**

Product name	Model
Endo Motor	Endo Radar
	Endo Smart
	E-Com、E-Com+
	EndoMatic
	Ai-Motor、MotoPex
	Endo Radar Plus
	Endo 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.: DD60137494 0001

Issue date: 2019-07-16

Expiry date: 2024-05-27

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

Company: **Guilin Woodpecker Medical Instrument Co., Ltd.**





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

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Address: Information Industrial Park, GuiLin National High-Tech Zone, GuiLin, GuangXi,  
541004, P.R.China

杨芸凤 2020.3.19  
Preparation, date

王淑萍 2020.3.19  
Review, date



Legally binding signature, Function

