

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

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

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+
	Endo Radar Pro
	Smart A
	Endo Pace、Endo Free



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

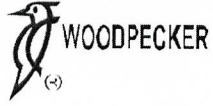
Certificate No.: DD60137494 0001

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



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

Company: Guilin Woodpecker Medical Instrument Co., Ltd.
Address: Information Industrial Park, GuiLin National High-Tech Zone, Guilin, GuangXi,
541004, P.R.China

苏婉婷 2022.1.5
Preparation, date

王政坤 2022.1.5
Review, date

于加康 2022.1.5
Legally binding signature, Function

