WOODPECKER

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

File No.: ZMN-YF-(R01)-01-001-02
Version: B

EC Declaration of Conformity

Manufacturer:

Guilin Woodpecker Medical Instrument Co., Ltd.

whose single Authorized Representative:

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

We, the manufacturer, herewith declare that the products
APEX LOCATORS UMDNS-*Code*:16355

Product name	Model
APEX LOCATOR	Woodpex I、Dpex I
	Woodpex III、Dpex III
	Minipex
	Woodpex V、DPEX V

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

杨芸凤 2020.3.19
Preparation, *date*

Review, *date* 2020.3.19

Legally binding signature, Function 2020.3.19

GUILIN WOODPECKER MEDICAL INSTRUMENT CO.,LTD 桂林市啄木鸟医疗器械有限公司 4503011007775