



EC Declaration of Conformity

Manufacturer:

whose single Authorized Representative:

Guilin Woodpecker Medical Instrument Co., Ltd.

**MedNet GmbH • Borkstrasse 10 • 48163
Muenster • Germany**

We, the manufacturer, herewith declare that the products,

Endo Motor, UMDNS-Code: 21400

Model:MotoPex

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 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 601151470001

Issue date: 2016-11-29

Expiry date: 2019-09-15

following the procedure relating to the EC Declaration of Conformity set out in Annex V 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

马清男 2019.6.27
Preparation , Date

王毅炜 2019.6.27
Review , Date

王毅炜 2019.6.27
Legally binding signature, Function

