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

EC Declaration of Conformity

Manufacturer:

Guilin Woodpecker Medical Instrument Co., Ltd.

whose single Authorized Representative:

MedNet GmbH • Borkstrasse 10 • 48163 Muenster • Germany

We, the manufacturer, herewith declare that
the products: *Piezo Bone Surgery* , UMDNS-Code: 18049

MODEL:Surgic Touch、Surgic Touch LED、Surgery-X、Surgery-X LED

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

苏琬婷 2018.12.10
Preparation , Date

2018.12.10
Review , Date

2018
Legally binding signature, Function