



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

File No.: ZMN/WI-07-067-02

Version: A

## 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

**Dental Diode Laser Device, GMDN-Code: 60340**

**MODEL: LX 16, LX 16 Plus, D-Laser 16, D-Laser blue**

meet the provisions of Directive 93/42/EEC which apply to them.

The medical device has been assigned to class IIb 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 II 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.: HD 2158053-1

Issue date: 2021-05-19

Expiry date: 2024-05-26

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

杨艺凤 2021.5.20  
Preparation, Date

王毅伟 2021.5.20  
Review, Date

Legally binding signature, Location, Function  
Guilin, PRRC

