

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



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

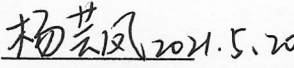
Certificate No.: HD 2158053-1
Issue date: 2021-05-19
Expiry date: 2024-05-26

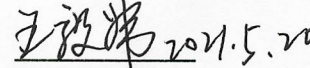
following the procedure relating to the EC Declaration of Conformity set out in Annex II of Directive 93/42/EEC.

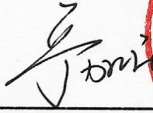
This Declaration of Conformity covers all medical devices as specified in the product list belonging to this declaration and is only valid in connection with a batch specific Certificate of Compliance for all products concerned bearing the CE mark

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


Preparation, Date


Review, Date


Legally binding signature, Location, Function

