

# EC Declaration of Conformity

*Manufacturer:*

*whose single Authorized Representative:*

Zhuolu Jontelaser Manufacturing Technology Co., MedNet GmbH  
Ltd.

No. 31, Sanguanmiao Alley, Zhuolu Town, Zhuolu County, Zhangjiakou City, Hebei  
Borkstrasse 10 . 48163 Muenster . Germany

We, the manufacturer, herewith declare that the products  
**Dermatological Diode Laser Systems**  
Model: T5PRO/T8PRO  
GMDN-Code: 58786

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

The medical device has been assigned to class IIb according to Rule 9, 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 (without II.4) of Directive 93/42/EEC.

Compliance of the designated product with the Directive 93/42/EEC has been assured via assessment of the quality management system by the Notified Body

**TÜV Rheinland LGA Products GmbH**  
**Tillystraße 2, 90431, Nürnberg, Germany**

Certificate No.:HD 60145283 0001

Issue date: 2020-02-17

Expiry date: 2024-5-26

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: Zhuolu Jontelaser Manufacturing Technology Co., Ltd.

Address: No. 31, Sanguanmiao Alley, Zhuolu Town, Zhuolu County, Zhangjiakou City, Hebei

2020.06.16

Place, date

Legally binding signature, Function



File no.: JS- T8/T5 PRO-0004

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