

File No.: ZMN-YF-(RD01)-01-005-02

Version:H

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, here with declare that the products **Ultrasonic Scalers**, *UMDNS-Code*: **17497**

Product name	Model
	D3、D3 LED
	D5、D5 LED
	D6、D6 LED
	D7、D7 LED
	D600、D600 LED
	S6、S6 LED
	U6、U6 LED
	U600、U600 LED
	UDS-A、UDS-A LED
	UDS-K、UDS-K LED
	UDS-E、UDS-E LED
	UDS-L、UDS-L LED
	UDS-P、UDS-P LED
	UDS-N2、UDS-N2 LED
	UDS-N3、UDS-N3 LED
	UDS-N6、UDS-N6 LED
	V2、V2 LED
	V3、V3 LED
	V6、V6 LED
Dental Scaler and Air Polisher	PT-A, AP-B
Dental Air Polisher	AP-A



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



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The product concerned has been manufactured under a quality management system according to Annex VI and Annex VII 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 60137494 0001 Issue date: 2019-7-16 Expiry date: 2024-5-27

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

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Review , Date . 12.1