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

File No.: ZMN/WI-07-048-02
Version: A.1

EC Declaration of Conformity

Manufacturer:

Guilin Woodpecker Medical Instrument Co., Ltd.

whose single Authorized Representative:

MedNet EC-REP GmbH•Borkstrasse 10 • 48163 Muenster • Germany

We, the manufacturer, herewith declare that the products
Gutta Percha Obturation Device, *GMDN-Code:* 44859
MODEL:Fi-P , Fi-G, Fi-E

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 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

陈燕 2022.3.9	王丽娟 2022.3.9	2022.3.9
Preparation , Date	*Review , Date*	*Legally binding signature, Location, Function*