

EC Declaration of Conformity

Manufacturer: Guilin Woodpecker Medical Instrument Co.,Ltd
Information Industrial Park, Guilin National High-Tech
Zone, Guilin, 541004 Guangxi, P.R. China

SRN of Manufacturer: CN-MF-000009139

European Representative: MedNet EC-REP C Iib GmbH
Borkstrasse 10, 48163 Muenster, Germany

SRN of European Representative: DE-AR-000011194

Product Name: Apex Locator

Models: Woodpex I, Dpex I, Woodpex III, Dpex III, Woodpex X,
DPEX X, Woodpex V, DPEX V, Ai-Pex

Classification: II a / Rule 9 (MDR Annex VIII)

EMDN Code: Z121101

Basic UDI-DI: 694484360080KE

Conformity Assessment Route: Annex IX Chapters I and III of (EU)2017/745, MDR

We, the manufacturer, herewith declare under our sole responsibility that the above mentioned products meet the provisions of the Regulation (EU) 2017/745 on Medical Devices (MDR) and other relevant Union legislation that provides for the issuing of an EU declaration of conformity. All supporting documentations are retained under the premises of the manufacturer.

Notified Body: TÜV Rheinland LGA Products GmbH
Tillystraße 2 · 90431 Nürnberg
Country: Germany

CE Certificate No.: NB Identification number: 0197
HZ 2158053-1

Effective date: 2023-07-31

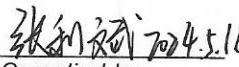
Expire date : 2028-07-30

Name: Ning Jiakang

Position: PRRC

Location: Guilin, China

Signature: 


Complied by:


Reviewed by: