

EU DECLARATION OF CONFORMITY

According to Art. 19 of Regulation (EU) 2017/745 on Medical Devices

Manufacturer: Guangzhou Nuoning Medical Supplies Co., Ltd
3rd Floor, Building B14, Huachuang Animation Industrial Park, No. 9 Huateng Road, Jinshan Village, Shiji Town, Panyu District, Guangzhou. 511450, China.

Trademark: N. A.

SRN: Not available yet

European Representative: MedPath GmbH
Mies-van-der-Rohe-Strasse 8
80807 Munich, Germany

SRN: DE-AR-000000087

Trade name: Venom Extractor

Product Name: Venom Extractor

Product code / Catalogue number

Basic UDI 6973448020002

Classification acc. to MDR Ax. VIII: Class I, rule 1

Applied Common Specifications / Standards: EN ISO 14971:2019+A11:2021
EN ISO 13485:2016+A11:2021

Conformity assessment procedure: N. A.

CE certificate No.: N. A.

Name and ID of the Notified Body: N. A.

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). All supporting documentations are retained under the premises of the manufacturer.

Li Nianwen,
General Manager

Li Nianwen

Guangzhou 16. 05. 2023

CE/MDR-MC-01-02