

## EU Declaration of Conformity

Manufacturer: **XUZHOU MAICUFF TECHNOLOGY CO.,LTD**

**8#-2-1106 Jinxiujiayuan, Jianguo West Road, Xuzhou City, China**

Factory Add: **3<sup>rd</sup> Floor, D Building, NO.47 Zhujiang Road, Tongshan District, Xuzhou City, China**

EC Representative: **MedNet EC-Rep GmbH**

Address: **Borkstrasse 10, 48163, Muenster, Germany**

Declares under our sole responsibility that the CE marked medical devices:

Product and Trade Name : **NIBP Cuff(Including NIBP Air Hose and Cuff Connector)**

Basic UDI-DI: **697133158M50017G, 697133158M50027J**

UMDNS Code: **11073**

Risk Class(**MDR EU 2017/745 Annex VIII, Chapter III, 4.1**): **Class I, Rule 1**

Conformity Assessment Route: **Annex II and Annex III**

A statement that the device that is covered by the present declaration is in conformity with the Medical Device Regulation/MDR (EU) 2017/745 and other relevant Union legislation.

Manufacturer is exclusively responsible for the DoC.

Place and Date: **Xuzhou City, May 18, 2021**

Name: **Li Zhaoqian**

Function: **General Manager**

Signature:

