DECLARATION OF CONFORMITY

ACCORDING TO (EU) 2017/745 MEDICAL DEVICE REGULATION

EU Representative

SUNGO Europe B.V.
Fascinatio Boulevard 522, Unit 1.7,
2909VA Capelle aan den IJssel, The Netherlands
SRN: NL-AR-000000247

Conformity Assessment

Conformity Assessment Procedure
Annex II+III of Regulation (EU) 2017/745

Applicable Standards
EN ISO 14971: 2019
EN ISO 15223-1: 2021
EN ISO 20417:2021
EN ISO 10993-1: 2020
EN ISO 10993-5: 2009
EN ISO 10993-10:2023
EN ISO 10993-23:2021
EN 60601-1-2:2015/A1:2021
EN 60601-1:2006/A2:2021
EN 12184: 2014
EN 62366-1:2015

Remark

The declaration of conformity is valid in connection with the release technical document CE/MDR-Y122127-06.

All the supporting documentation is retained at the premises of the manufacturer.

The Declaration of Conformity is exclusively under the sole responsibility of the manufacturer.

Manufacturer

Name: Zhongshan A&J Medical Equipment Co., Ltd.
Address: L3 Shenghui South Road, Nantou Town, Zhongshan City, Guangdong, 528427 China.
SRN:

Product Information

Name: Electric Wheelchair

Model: Viper; Viper Plus; Viper Lift
EMDN: Y122127
Basic UDI: 697786346EwheelchairEK
Classification: Class I, According to Rule 13, Annex VIII, Regulation (EU) 2017/745

Declaration

We herewith declare that the above-mentioned products meet the requirements of Medical Device Regulation (EU) 2017/745 and the applicable standards above.

Signature: Date: 2023.8.30

Position: GM Place: Guangdong/China