

# EU DECLARATION OF CONFORMITY

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

**Manufacturer:** Suzhou Sweetrich Vehicle Industry Technology Co., Ltd.  
Building B, No.7 Chao Qian Road, Suzhou Industrial Park , Suzhou, Jiangsu Province, China  
Zip code:215000

**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:** **Mobility Scooter**

**Product Name:** **Mobility Scooter**

**Models:** SW1000A,SW1000S,SW1000L,SW100SL,SW1000F,SW1000S PLUS,S1Lite,Max Plus,Max  
Sport,JD001,JD002,SW1250,SW1200,SW1260,SW1400, SW1400D,SW1700,SW1800,SW3100,SW3200,SW3300,SW1000M, SW1000A2,SW1000A3,1000N,1001N,1000D.

**Basic UDI:** 697307988SWSCOOTERXN

**Classification acc. to MDR Ax. VIII:** Class I, rule 13

**Applied Standard & Common Specification:** EN 12184:2014、 ISO 14971: 2020

**Conformity assessment procedure:** Annex II + Annex III of MDR

**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.

Signature of issue person:

Suzhou, 02. 12. 2024

Name: Ryan



Position:General Manager