EU DECLARATION OF CONFORMITY

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

Manufacturer:	Suzhou Sweetrich Vehicle Industry Technology Co., Ltd. No.68 Xinfa Road,Loufeng Town,Suzhou Industrial Park,Suzhou City,Jiangsu Province,China Zip code:215123
SRN:	CN-MF-000011210
European Representative:	MedPath GmbH Mies-van-der-Rohe-Strasse 8 80807 Munich, Germany
SRN:	DE-AR-000000087
UK Responsible Person:	MedPath Limited 27 Old Gloucester Street, London, WC1N 3AX United Kingdom
Trade name:	**Mobility Scooter**
Product name:	**Mobility Scooter**
Model Number:	SW1000A,SW1000A-1,SW1000E,S1 Sport,SW1000L, SW1000F, SW1000SL, S1 Plus, S1 Lite, Max Plus, Max Sport, JD001, JD002, S20, S25, S28, S70, S50(SW1200), SW1400D, SW3200, SW3300
Basic UDI:	697307988SWSCOOTERXN
Classification acc. to MDR Ax. VIII:	Class I, rule 13
Applied Standard & Common Specification:	EN 12184:2014
Conformity assessment procedure:	MDR Article 19 + Annex II + Annex III
CE certificate No.:	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,01.02 . 2023

Name:Ryan

Position:General Manager