



DECLARATION OF CONFORMITY

Regarding UK Medical Devices Regulations 2002

Manufacturer: Wiseton Industries Limited

Address: No.9 Jiahe Road, Jiashan County, Jiaxing City, Zhejiang Province, P.R. China

UK Responsible Person: SUNGO Certification Company Limited

Address: 3rd floor, 70 Gracechurch Street, London. EC3V 0HR

Product Name: Electric Patient Lift

Model: PL300, PL301, PL350

Classification: Class I

Conformity Assessment Procedure: Rule 12, Annex IX, Directive 93/42, Part II of the UK MDR 2002

We herewith declare that the above-mentioned products meet the requirements of UK Medical Devices Regulations 2002 and the following standards.

BS EN ISO 20417:2021 BS EN ISO 14971:2019 BS EN ISO 15223-1:2021
BS EN ISO 10993-5:2009 BS EN ISO 10993-1: 2020 BS EN ISO 10993-10: 2023
BS EN ISO 10993-23:2021 BS EN 60601-1:2006+A13:2024
BS EN 60601-1-2:2015 + A1:2021

Signature: 

Position: GM

Date: 2025-04-14

Place: Zhejiang/ China



The declaration of conformity is valid in connection with the release technical document UKCA-TCF-001.

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