



# 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** Rule 12, Annex IX, Directive 93/42, Part II of the UK MDR  
**Procedure:** 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*