



# DECLARATION OF CONFORMITY

ACCORDING TO (EU) 2017/745 MEDICAL DEVICE REGULATION

## EU Representative

SUNGO Europe B.V.  
Olympisch Stadion 24, 1076DE  
Amsterdam, 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: 2016  
EN 1041:2008+A1:2013  
EN ISO 10993-1: 2020  
EN ISO 10993-5: 2009  
EN 12184:2014

### Remark

*The declaration of conformity is valid in connection with the release technical document*

*CE/MDR-EMD-01.*

*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:** Zhejiang Innuovo Rehabilitation Devices Co., Ltd

**Address:** No.196 Industry Road, Hengdian Movie Zone, Dongyang, Zhejiang, China

**SRN:** CN-MF-000008727

## Product Information

**Name:** Power Wheelchair

**Model:** W5521(W5521-SIL) (#IFPC17SIL),  
W5517(W5517-BLK) (#AFPC17BLK)

**GMDN:** 40840

**Basic UDI-DI:** 697076597PW001QM

**Intended purpose:** The product is intended to provide transportation for disabled or elderly persons limited to a seated position.

**Classification:** Class I, According to Rule 1, 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: *Leo Zhou* Date: *Sept. 12, 2021*

Position: GM

Place: Zhejiang/China

浙江英诺华康复器材有限公司  
ZHEJIANG INNUOVO  
REHABILITATION DEVICES CO., LTD