CE Declaration of Conformity (MDD)

Under sole responsibility, the undersigned certify that the medical product(s) described hereafter as;
Product Name/Designation: Invacare DECPAC Ramp
Model(s): EBL/SEN/MP/PER

With the following locations;

Manufacturer: Shanghai Cedar Composites Technology Co., ltd
Address: No. 509 Haoge Road, Nicheng Town
District: Pudong New District, Shanghai
Country: China

Is (are) in conformity with;
Medical Device Directive 93/42/EEC-Annex VII_ as classification I, Article 4 of the RoHS Directive 2011/65/EU and Directive (EU) 2015/863 of the European parliament and the Council of 8 June 2011 for restriction of the use of certain hazardous substances in electrical and electronic equipment,

The following harmonized standard(s),
EN ISO 13485:2016
And using a quality management system certified to ISO 13485:2016 by Ingeer Certification Assessment Co, Certificate Number: 117 19 Q0M 0063 R1M

With medical Device Directive 93/42/EEC monitoring and supervision by DNV Business Assurance, as Notified Body 0434, Certificate Number: 83877-2010-CE-RGC-NA, Rev.2 to Article 11.3a and Annex II (excluding Section 4).

Signed by: Date: 5/25/2021 On behalf of : Shanghai Cedar

Name: Haiyan Wang Title: Plant Quality Manager