MD
CE

HOVERTECH International

EU DECLARATION OF CONFORMITY

Manufacturer	HoverTech International 4482 Innovation Way Allentown, PA 18109
SRN EC REP	US-MF-000008435
Authorized Representative Name	CEpartner4U
Authorized Representative Address	Esdoornlaan 13, 3951DB Maarn, The Nederlands
Statement	This EU declaration of conformity is issued under the sole responsibility of the manufacturer. The device(s) covered by present declaration is/are in conformity with EU Regulation 2017/745 on medical devices.
Basic UDI-DI	081629901HJSF
Intended purpose	In the event of a patient fall, the HoverJack® Air Patient Lift is used to lift the patient in a supine position from the floor to bed or stretcher height, utilizing the HoverTech Air Supply to inflate each of the four chambers.
Product / device name	HoverJack Evacuation HoverJack EMS Evacuation HoverJack
Risk class of the device	Class 1, rule 1
Place	Allentown, PA USA
Date of issue	January 30, 2023
Name and function	Susan Pavelko /Quality Manager

Susan Pavelko

Signature, on behalf of HoverTech International