MD
CE

HOVERTECH International

EU DECLARATION OF CONFORMITY

Manufacturer	HoverTech International 4482 Innovation Way Allentown, PA 18109
SRN	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	**081629901ASSC**
Intended purpose	The Air Supply is used along with HoverTech's Air assisted devices to assist caregivers with patient transfers, positioning, turning, and proning.
Product / device name	HT-AIR 2300
Risk class of the device	Class 1, rule 13
Place	Allentown, PA USA
Date of issue	19, January 2023
Name and function	Susan Pavelko /Quality Manager

Susan Pavelko

Signature, on behalf of HoverTech International