permobil

EU-declaration of conformity

As manufacturer
Permobil AB
Box 120
Per Uddéns väg 20
861 23 Timrå
Sweden

declares under the sole responsibility that below specified product is in conformity with the

Regulation (EU) 2017/745
on medical devices (MDR)

General description	Electrically powered wheelchair. Intended use: The intended use for F5 Corpus VS is to provide outdoor and indoor mobility, including a standup feature, to persons limited to a seated position that are capable of operating a powered wheelchair. The trade name of the wheelchair is **F5 Corpus VS** and is manufactured by Permobil AB. The manufacturer's internal article number is 108400-99-0.
Basic UDI-DI	**7330818COMDM**
SRN	SRN will be assigned when Eudamed is up and running.
Product class:	Class I (according to Annex VIII Chapter III, rule 1 and rule 13)
Conformity procedure used:	Annex IX
Standards	Conformity to the general safety and performance requirements have been demonstrated by using the following standards: EN ISO 13485 EN 12184 IEC 62304 EN ISO 14971 EN 1041 EN ISO 10993-1
Original drawn up	2020-03-17
Place and date:	Timrå, Sweden 2020-09-07

On behalf of Permobil AB

Jonas Lindström
Executive Vice President Business Unit Power Products, Permobil AB