

EU-declaration of conformity

As manufacturer

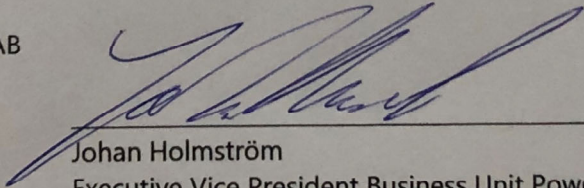
Permobil AB
Box 120
Per Uddéns väg 20
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Sweden

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 of the M3 Corpus powered wheelchair is to provide indoor and outdoor mobility to persons limited to a seating position that are capable of operating a powered wheelchair. The trade name of the wheelchair is M3 Corpus and is manufactured by Permobil AB. The manufacturer's internal article number is 107724-99-0.
Basic UDI-DI	7330818MXUF
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:	Göteborg, Sweden 2020-03-17

On behalf of Permobil AB



Johan Holmström

Executive Vice President Business Unit Power Products, Permobil AB