EU Declaration of Conformity

As manufacturer

Permobil AB
Klökanvägen 16
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Sweden

declares under the sole responsibility that below specified product is in conformity with the following relevant Union harmonization legislation:

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

Radio Equipment **Directive 2014/53/EU** (RED)

General description	Electrically powered wheelchair. Intended use: The intended use of the **F5 Corpus VS** powered wheelchair 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 108851-99-0.
Basic UDI-DI	**7330818COMDM**
SRN	SE-MF-000014317
Product class acc to MDR:	Class I (according to Annex VIII Chapter III, rule 1 and rule 13)
Standards (RED):	Conformity to the essential requirements of RED has been demonstrated by using the following standards: ISO 7176-9:2009 — EN 55032:2015 ISO 7176-14:2008 — EN 62311:2008 ISO 7176-21:2009 — ISO 14971:2019 Draft ETSI EN 301 489-19 v2.2.0 2020-09 Draft ETSI EN 301 489-52 v1.1.2 2020-12 EN 301 489-17 v3.2.4 2020-09 ETSI EN 300 328 v2.2.2 2019-07 ETSI EN 303 413 v1.1.1 2017-06 ETSI EN 301 511 v12.5.1 2017-03 ETSI EN 301 908 v13.1.1 2019-11 ETSI TS 151010-1 v13.11.0 Test case 12.2.1 and 12.2.2 2020-02
Original drawn up	2022-12-23
Place and date:	Sundsvall, Sweden 2024-09-02

On behalf of Permobil AB

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