

## EU Declaration of Conformity

As manufacturer  
**Permobil AB**  
Klökänvägen 16  
863 41 Sundsvall  
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 **M5 Corpus** powered wheelchair is to provide outdoor and indoor mobility to persons limited to a seated position that are capable of operating a powered wheelchair.  
The trade name of the wheelchair is **M5 Corpus** and is manufactured by Permobil AB. The manufacturer's internal article number is 109046-99-0.

**Basic UDI-DI:**

**7330818COMDM**

**SRN:**

SE-MF-000014317

**Product class according 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  
ETSI 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

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On behalf of Permobil AB

  
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