

**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 of the F3 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 **F3 Corpus** and is manufactured by Permobil AB. The manufacturer's internal article numbers are 108398-99-0.

**Basic UDI-DI**                      7330818FXTS

**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)

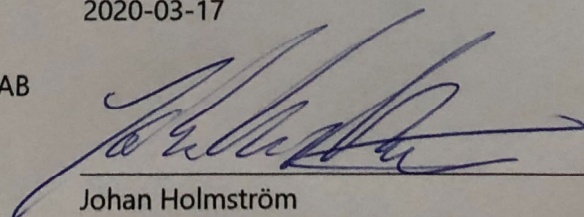
**Conformity procedure used:**      Annex IX

**Standards**                              Conformity to the general safety and performance requirements have been demonstrated by using the following standards:  
EN ISO13485  
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

  
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Johan Holmström  
Executive Vice President Business Unit Power Products, Permobil AB