permobil

EU Declaration of Conformity

CE

We hereby declare that the below-mentioned medical device meets the provisions of **Regulation (EU) 2017/745 of the European Parliament and of the Council** on medical devices (MDR).

Manufacturer	Permobil Progeo Active Design s.r.l. Vicolo Luigi Negrelli, 5 31038 Paese Italy
General description	MOTOTRONIK Intended use: **MOTOTRONIK** is an electric handbike designed to be used with manual wheelchairs, in order to provide motorized propulsion, improving user mobility and independence. **Target users:** • Adults and adolescents who are trained, informed, and aware of the associated risks. • Users must have sufficient visual, physical, and cognitive abilities to ensure safe use. **Restrictions:** • Not intended for children or for individuals who are untrained or physically unfit. **Device characteristics:** • Made of aluminum, steel, plastic, and rubber/PU. • Operates only in contact with intact skin. The commercial name of the medical device is **MOTOTRONIK**. The trade name of the medical device is **MOTOTRONIK** The manufacturer's internal article number(s) is/are **MK**
Basic UDI-DI	**8033172DEVPOWER3S**
SRN	**IT-MF- 000051528**
Device classification	Classe I (secondo Allegato VIII, Capitolo III, regola 1)
First CE marked	2021-09-01
Place and date	Paese, Italia

This EU Declaration of Conformity is issued under the sole responsibility of the Manufacturer.

On behalf of
Permobil Progeo Active Design S.r.l. 2026-01-23

Giulia Guidolin
Site Quality Manager Italy Giulia Guidolin