EU Declaration of Conformity

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	**DUKE** Intended use: Progeo DUKE is an ultra-light, rigid carbon fiber wheelchair for self-propulsion. Designed for daily use by both active users and those with severe mobility limitations, it ensures safe indoor and outdoor movement within -30°C to +60°C. Caregiver assistance is recommended for users unable to move independently. Not suitable for rough terrain, steep slopes, or saline environments. The trade name of the medical device is **DUKE** The manufacturer's internal article number(s) is/are **DK**
Basic UDI-DI	**8033172DEVWHEELZD**
SRN	**IT-MF-000051528**
Device classification	Class I (according to Annex VIII Chapter III, rule 1)
First CE marked	2017-06-01
Place and date	Paese, Italy 2026-01-23

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

On behalf of Permobil Progeo Active Design s.r.l.

Giulia Guidolin
Site Quality Manager Italy