EF-0938 – UKCA Declaration of Conformity

Version 2

Product name	**Invacare Propad Cushion Range** **PU Foam Cushion (PL3655, PL3792)**
Basic UDI-DI	**505505PROPADANDPROFLEXD6**
Intended purpose	**The cushion is to be used in an armchair or wheelchair as part of an overall pressure injury prevention program of care. Propad RevolveV can be used for "High Risk" users. Propad® RevolveSi can be used for "Very High Risk" users.**
Manufactured by	**Invacare UK Operations Ltd,** **Pencoed Technology Park,** **Pencoed Bridgend, CF35 5AQ**
Manufacturer Single Registration Number (SRN)	**GB-MF-000007111**
Authorized Representative (EU-REP)	**Invacare (Portugal), Unipessoal, Lda.** **Rua Estrada Velha, 949** **4465-784 Leca do Balio** **Portugal**
Authorized Representative SRN	**PT-AR-000006129**
United Kingdom Responsible Person (UKRP)	**N/A**

This UKCA declaration of conformity is issued under the sole responsibility of the manufacturer.

Device Classification according to Directive 93/42/EC, Annex IX	**Class I**
The object of the declaration is in conformity with the relevant legislation:	**UK MDR 2002 (As amended)** **REACH 1907/2006,** **RoHS 2011/65/EU,**
Applied harmonised standards, common specifications, national standards or other normative documents	**EN1021:2014 – Part 1 and 2** **BS EN ISO1497:2019/A11-2021** **ISO 10993-5:2009** **ISO 9999:2022** **ISO 21856:2022**
Initial date of first Declaration of Conformity	**1985**

Place and issue date:

Pencoed, 13-Jan-2026

Operations Manager

Darren Swetman,