

EF-0938 – UKCA Declaration of Conformity

Version 2



Yes, you can.

Product name	Invacare Propad Cushion Range (Propad Premier Cushion, Propad Original Cushion, Propad Profile Cushion, Propad Revolve Si Cushion, Propad Revolve V Cushion.)
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, 07-Oct-2026

Regulatory Affairs Manager

Renee de Beet,

Electronically signed by: Renee de Beet
Reason: Reviewer/Approver - I have reviewed and approve this document.
Date: 07-Oct-2026 11:02:26 GMT+1