

EF-0938 – UKCA Declaration of Conformity

Version 2



Product name	Invacare Matrix Flo-tech Cushion Range Moulded PU foam cushion with or without gel sack (PL3650, PL3651, PL3652, PL3653, PL3654, PL3656, PL3657, PL3658, PL3659, PL3661, PL3665, PL3666, PL5346)
Basic UDI-DI	505505MATTRXFLOTECHT3
Intended purpose	These cushions are intended to be used in an armchair or wheelchair as part of an overall pressure injury prevention program of care
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	1995

Place and issue date:

Operations Manager

Pencoed, 13-Jan-2026

Darren Swetman,