Document Reference: Document Issue Number: Document Issue Date: Page 1 of 2 ME-EUDOC-034 01 23/06/2022



EU Declaration of Conformity

This European Declaration of Conformity is issued under the sole responsibility of the manufacturer.

MANUFACTURER					
Name of Company	Registered Office	Trading Address	SRN		
Medicare Products Ltd	South Fen Road, Bourne, Lincolnshire, PE10 0DN. UK	Unit B, Dolphin Way, Purfleet, Essex, RM19 1NZ, UK	GB-MF-000011612		

AUTHORIZED REPRESENTATIVE					
Name of Company	Address	SRN	Phone/email		
International Associates	The Black Church, St		+353 16971561		
Auditing & Certification	Mary's Place, Dublin 7,	IE-AR-000002248	EUAR@ie.ia-net.com		
Limited	D07 P4AX Ireland		EOAK@ie.ia-fiet.com		

Product Name	Code / Catalog Number	Basic UDI-DI
Nitrile Powder Free Sterile Cleanroom Gloves (NITREX 600)	GN38	N/A
Intended Purpose	Photo	
Cat III PPE for use in cleanrooms and offering a degree of chemical protection against the chemicals specified in the IFU.		

RISK CLASS FOR MEDICAL DEVICES				
Device Classification Common Specifications / Standards		Common Specifications / Standards		
Class:	N/A	N/A		
Rule:	N/A	IN/A		

Medicare Products Ltd declares that the above-mentioned product meets the provision of the following EU legislation:

- Is classed as Cat III PPE in conformity with the provisions of Regulation (EU) 2016/425 on personal protective equipment and with National Standards transposing harmonised standards EN 374-1:2016, EN 374-5:2016, and EN 420:2003+A1:2009.
- Is identical to the personal protective equipment which is the subject of EU Type-Examination Certificate number 2777/11793-01/E05-01, issued by SATRA Technology Europe Limited, Bracetown

T: +44 (0) 1708 863868 E: enquiries@medicareproducts.com W. www.medicareproducts.com

Document Reference: Document Issue Number: Document Issue Date: Page 2 of 2 ME-EUDOC-034 01 23/06/2022



Business Park, Clonee, Dublin, D15 YN2P, Republic of Ireland (Notified Body number 2777), according to Annex V (Module B) of Regulation (EU) 2016/425.

• Is subject to the quality assurance conformity of the production process Module D set out in Annex VIII of Regulation (EU) 2016/425, under the surveillance of the notified body SGS FIMKO OY, Särkiniementie 3, 00211 Helsinki, Finland (Notified Body Number 0598).

COMPANY REPRESENTATIVE: David Langridge

TITLE: Head of Technical SIGNATURE: Designing

PLACE: Bourne, UK DATE: 23 June 2022