Document Reference:PH-DOC-079Document Issue Number:07Document Issue Date:20/02/2024Page 1 of 220/02/2024



EU Declaration of Conformity

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

MANUFACTURER				
Name of Company	Address	SRN		
Polyco Healthline Ltd	South Fen Road, Bourne, Lincolnshire, PE10 0DN. UK	GB-MF-000015289		

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

EU IMPORTER	
Name of Company	Address
Polyco Healthline BV	7 th Floor, De Boelelaan 7, Amsterdam, 1083HJ

PRODUCT IDENTIFICATION			
Product Name	Code / Catalogue Number	Basic UDI-DI	
Blue Nitrile Powder Free Examination Glove	GN91	5024951GNPFNS00YA	
Intended Purpose	Photo		
Examination gloves for non-invasive use to prevent contact with skin, bodily fluids, and chemicals. Invasive with respect to natural body orifices.	N	4	

RISK CLASS FOR MEDICAL DEVICES			
Device Classification Common Specifications / Standards		ications / Standards	
Class:	I	EN455-1:2020 EN455-2:2015	Medical gloves for single use: Freedom from holes Medical gloves for single use: Physical properties
Rule:	5	EN455-3:2015 EN455-4:2009	Medical gloves for single use: Biological evaluation Medical gloves for single use: Shelf-life determination

RISK CATEGORY OF PERSONAL PROTECTIVE EQUIPMENT AND PERFORMANCE LEVELS

Product Risk Category			
Personal Protective Equipment (PPE) Category III			
Standard	Performance Levels		
EN ISO 374-1:2016+A1:2018	Туре С - К		
EN ISO 374-5:2016	Protection against bacteria and fungi	Protection against viruses	
	Pass	Pass	

Head Office & Distribution Centres: South Fen Road, Bourne, Lincolnshire, PE10 0DN, UK

Document Reference:PH-DOC-079Document Issue Number:07Document Issue Date:20/02/2024Page 2 of 2



Polyco Healthline Ltd declares that the above-mentioned products meet the provision of the following EU legislation:

- Medical Devices Regulation (EU) 2017/745
- 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 ISO 374-1:2016+A1:2018, EN ISO 374-5:2016, and EN ISO 21420:2020.
- Is identical to the personal protective equipment which is the subject of EU Type-Examination Certificate number 2777/11509-06/E00-00, issued by SATRA Technology Europe Limited, Bracetown 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 conformity assessment procedure Module C2 set out in Annex VII of Regulation (EU) 2016/425, under the surveillance of the notified body SATRA Technology Europe Limited, Bracetown Business Park, Clonee, Dublin, D15 YN2P, Republic of Ireland (Notified Body number 2777).

COMPANY REPRESENTATIVE:	David Langridge		
TITLE:	Head of Technical	SIGNATURE:	D Longridge
PLACE:	Bourne, UK	ISSUE DATE:	20 th February 2024