

Innovation & Quality

EU DECLARATION OF CONFORMITY

Manufacturer : Hartalega NGC Sdn. Bhd.

Manufacturer's Address : No.1, Persiaran Tanjung, Kawasan Perindustrian Tanjung,

43900 Sepang, Selangor Darul Ehsan, Malaysia.

EU Representative : MDSS GmbH

Schiffgraben 41, 30175 Hannover, Germany. Reference Only

Product Description (EU MDR) : Nitrile Powder Free Examination Gloves

Device Classification (EU MDR) : Class I, according to Annex VIII of Regulation (EU) 2017/745

Rule(s) : 1 and 5

Conformity Assessment : Annex II and Annex III

Procedure

Basic UDI-DI : 955100777HNGCTFMD005AQ8

Authorized Representative SRN : DE-AR-000005430

Manufacturer SRN For Refereing MY-MF-000010459 eference Only

Product Description (EU PPER) : _ Nitrile Powder Free Examination Gloves (≥ 2.2 mil)

Available in standard minimum 240mm length or a longer Communication of the length of

cuff variant of 280mm

Device Classification (EU PPER) : Category III (Type B)

EU Type-Examination : 2777/11578-04/E00-00

Certificate Number (EU PPER)

For Ref Intended Purpose For Refere: Nitrile Powder Free Examination Gloves are intended to be

used to contribute to prevent cross contaminations in the framework of medical examinations and diagnostic/therapeutic procedures conducted under non-sterile

conditions.

and

Nitrile Powder Free Examination Gloves is intended to protect users from substances and mixtures which are

hazardous to health and harmful biological agents that may cause very serious consequences or irreversible damage to

health.

Standard Reference : Attachment I

Reference to Trade Name : Attachment II

Hartalega Holdings Berhad (741883-X)

C-G-9, Jalan Dataran SD1, Dataran SD PJU 9 Bandar Sri Damansara 52200 Kuala Lumpur, Malaysia Tel: +603 - 6277 1733 Fax: +603 - 6280 2533 www.hartalega.com.my Hartalega NGC Sdn Bhd (984586-P)

No. 1, Persiaran Tanjung Kawasan Perindustrian Tanjung PREDICE 43900 Sepang, Selangor, Malaysia Tel: +603 - 8707 3000 Rev 33

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We, Hartalega NGC Sdn. Bhd. herewith declared that above mentioned device:

- The Council of medical devices.
 - is in conformity with the provisions of Regulation (EU) 2016/425 on personal protective equipment.
 - is subject to the conformity assessment procedure Module D set out in Annex VIII of Regulation (EU) 2016/425, under the surveillance of the notified body SATRA Technology Europe Limited, Bracetown Business Park, Clonee D15YN2P, Republic of Ireland (Notified Body number 2777).

This EU declaration of conformity is issued under the sole responsibility of the manufacturer, Hartalega NGC Sdn. Bhd.

Place and Date of Issue : Hartalega NGC Sdn. Bhd./ 07th May 2025

Signed on Behalf of Hartalega NGC

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Position: GENERAL MANAGER - QUALITY
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Standard	han Only For Rate Title Only For Rater ance Or
ISO 9001:2015	Quality Management Systems – Requirements
EN ISO 13485:2016+A11:2021	Medical Devices – Quality Management Systems – Requirements for Regulatory Purposes
EN 455-1:2020+A2:2024	Medical Gloves for Single Use Part 1: Requirements and Testing for Freedom from Holes
3S EN 455-1:2020+A2:2024	Medical Gloves for Single Use Part 1: Requirements and Testing for Freedom from Holes
EN 455-2:2024	Medical Gloves for Single Use Part 2: Requirements and Testing for Physical Properties
BS EN 455-2:2024	Medical Gloves for Single Use Part 2: Requirements and Testing for Physical Properties
EN 455-3:2023	Medical Gloves for Single Use Part 3: Requirements and Testing for Biological Evaluation
BS EN 455-3:2023	Medical Gloves for Single Use Part 3: Requirements and Testing for Biological Evaluation
EN 455-4:2009 nce Only	Medical Gloves for Single Use Part 4: Requirements and Testing for Shelf Life noe Only Determination
nce Only For Refere BS EN 455-4:2009 For Reference Only	Medical Gloves for Single Use The Control of Part 4: Requirements and Testing for Shelf Life Determination
BS EN ISO 20417:2021	Medical devices – Information to be Supplied by the Manufacturer
BS EN ISO 14971:2019+A11:2021	Medical Devices – Application of Risk Management to Medical Devices – For Reference Only
ISO 15223-1:2021 or Refere	Medical Devices –Symbol to be Used with Information to be Supplied by the Manufacturer — Conty — For Reference Or Part 1: General Requirements
For Reference Only ISO 10993-1:2018 Ince Only For Refere	Biological Evaluation of Medical Devices Part 1: Evaluation and Testing within a Risk Management Process
BS EN ISO 10993 – 1: 2020	Biological Evaluation of Medical Devices Part 1: Evaluation and Testing within a Risk Management Process
ISO 10993-5:2009 OF Refere	Biological Evaluation of Medical Devices Ty For Reference Or Part 5: Tests for In Vitro Cytotoxicity
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Standard	Title	
BS EN ISO 10993 – 5:2009	Biological Evaluation of Medical Devices Ference Only Part 5: Tests for In Vitro Cytotoxicity	
ISO 10993-10:2021	Biological Evaluation of Medical Devices Part 10: Tests for Skin Sensitization	
BS EN ISO 10993 - 10:2023	Biological Evaluation of Medical Devices Part 10: Tests for Skin Sensitization	
ISO 10993-11:2017	Biological Evaluation of Medical Devices Part 11: Tests for Systemic Toxicity	
BS EN ISO 10993 - 11:2018	Biological Evaluation of Medical Devices Part 11: Tests for Systemic Toxicity	
ISO 10993-18:2020/ Amd 1:2022	Biological Evaluation of Medical Devices Part 18: Chemical Characterization of Medical Device Materials Within A Risk Management Process Amendment 1: Determination of the uncertainty factor	
BS EN ISO 10993 – 18:2020+A1:2023	Biological Evaluation of Medical Devices Part 18: Chemical Characterization of Medical Device Materials Within A Risk Management Process	
ISO 10993-23:2021 Refere	Biological Evaluation of Medical Devices Part 23: Tests for Irritation	
BS EN ISO 10993 – 23:2021	Biological Evaluation of Medical Devices Part 23: Tests for Irritation	
ISO 2859-1:1999/Amd1:2011	Sampling Procedures for Inspection by Attributes Part 1: Sampling Schemes Indexed by Acceptance Quality Limit (AQL) for Lot-By-Lot Inspection	
ASTM D4169-23e1	Standard Practice for Performance Testing of Shipping Containers and Systems	

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Standard	For Reference Only Title or Reference Only	For Refer
EN ISO 21420:2020	Protective gloves - General requirements and test methods	anaa Only
EN ISO 374-1:2016+A1:2018	Protective gloves against dangerous chemicals and micro- organisms - Part 1: Terminology and performance requirements for chemical risks	For Refer
EN ISO 374-5:2016	Protective gloves against dangerous chemicals and micro- organisms - Part 5: Terminology and performance requirements for micro- organisms risks	ence Only For Refer
EN 421:2010 For Refer	Protective gloves against ionizing radiation and radioactive contamination	ence Only

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For Refere	Vasco Nitril blue	M: 9209825		
nce Only	For Reference Only For Refer	xL: 9209841 ence Only		
For Refere	nce Only For Reference Only Vasco Nitril white (150pcs)	For Refere S: 9208402 S: 9208410 M: 9208429		
nce Only	For Reference Only For Reference			
For Refere		S: 9207910		
nce Only	Vasco Nitril white (100pcs) For Reference Only For Reference	M: 9207929 L: 9207937 rence Only XL: 9207945		
For Refere	nce Only For Reference Only			
nce Only	For Ref Vasco Nitril Soft white or Refere	XL: 9201140		
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