

## **EU Declaration of Conformity**

Manufacturer : Hartalega NGC Sdn. Bhd.

Manufacturer's Address : No.1, Persiaran Tanjung, Kawasan Perindustrian Tanjung, 43900

Sepang, Selangor, Malaysia.

EU Representative : Medical Device Safety Service (MDSS)

Schiffgraben 41, 30175 Hannover, Germany.

Product Description (MDR) : Nitrile Powder Free Examination Gloves

Intended Purpose (MDR) : 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.

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

Rule (s) : 1 and 5

**Conformity Assessment** 

Procedure

: Annex II and Annex III

Basic UDI-DI : 955100777HNGCTFMD005AQ8

**Authorised Representative** 

SRN

DE-AR-000005430

Manufacturer SRN : MY-MF-000010459

Reference to Trade Name

(MDR)

Attachment I

Standard Reference (MDR) : Attachment II

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

Available in standard minimum 240mm length or a longer cuff

variant of 280mm

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

EU Type-Examination

Certificate Number (PPER)

: 2777/11578-02/E00-00

Reference to Trade Name

(PPER)

Attachment III

Standard Reference (PPER) : EN 420:2003+A1:2009

EN ISO 374-1:2016+A1:2018

EN ISO 374-5:2016

Hartalega Holdings Berhad (741883-X)

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



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

- is in conformity with the Regulation (EU) 2017/745 of The European Parliament and of 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 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 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./ 17th September 2021

Signed for and on Behalf of Hartalega NGC : Sdn. Bhd.

Name : NURUL AISYAH KONG

Position: DEPUTY GENERAL MANAGER -

**QUALITY ASSURANCE** 

## ATTACHMENT I

Product or Trade Name	Reference Number
Peppler Nitril Cloud 436	XS: 436XS S: 436S M: 436M L: 436L XL: 436XL

## **ATTACHMENT II**

Standard	Title
ISO 9001:2015	Quality Management Systems – Requirements
EN ISO 13485:2016	Medical Devices – Quality Management Systems – Requirements for Regulatory Purposes
EN 455-1:2000	Medical Gloves for Single Use Part 1: Requirements and Testing for Freedom from Holes
EN 455-1:2020	Medical Gloves for Single Use Part 1: Requirements and Testing for Freedom from Holes
EN 455-2:2015	Medical Gloves for Single Use Part 2: Requirements and Testing for Physical Properties
EN 455-3:2015	Medical Gloves for Single Use Part 3: Requirements and Testing for Biological Evaluation
EN 455-4:2009	Medical Gloves for Single Use Part 4: Requirements and Testing for Shelf Life Determination
BS EN 1041:2008+A1:2013	Information Supplied by the Manufacturer of Medical Devices
BS EN ISO 14971:2019	Medical Devices – Application of Risk Management to Medical Devices
ISO 15223-1:2016	Medical Devices – Symbols to be Used with Medical Device Labels, Labelling and Information to be Supplied Part 1: General Requirements
ISO 10993-1:2018	Biological Evaluation of Medical Devices Part 1: Evaluation and Testing within a Risk Management Process
ISO 10993-5:2009	Biological Evaluation of Medical Devices Part 5: Tests for In Vitro Cytotoxicity
ISO 10993-10:2010	Biological Evaluation of Medical Devices Part 10: Tests for Irritation and Skin Sensitization
ISO 10993-11:2017	Biological Evaluation of Medical Devices Part 11: Tests for Systemic Toxicity
ISO 10993-18:2005	Biological Evaluation of Medical Devices Part 18: Chemical Characterization of Materials
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

## ATTACHMENT III

Product or Trade Name	Reference Number
Peppler Nitril Cloud 436	XS: 436XS S: 436S M: 436M
	L: 436L XL: 436XL