

Innovation & Quality

EU Declaration of Conformity

Manufacturer	:	Hartalega Sdn. Bhd.	
Manufacturer's Address	:	C-G-9, Jalan Dataran SD1, Dataran SD PJU9, Bandar Sri Damansara, 52200 Kuala Lumpur, Malaysia.	
EU Representative	:	Medical Device Safety Service (MDSS) Schiffgraben 41, 30175 Hannover, Germany.	
Product Description (MDR)	:	Latex Powder Free Examination Gloves	
Intended Purpose (MDR)	:	Latex 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	:	955524480HSBTFMD001A57	
Authorised Representative SRN	:	DE-AR-000005430	
Manufacturer SRN	:	MY-MF-000010461	
Reference to Trade Name (MDR)	i	Attachment I	
Standard Reference (MDR)	:	Attachment II	
Product Description (PPER)	C	Latex Powder Free Examination Glove	
Device Classification (PPER)		Category III (Type B)	
EU Type-Examination Certificate Number (PPER)	:	2777/11080-03/E00-00	
Reference to Trade Name (PPER)	÷	Attachment III	
Standard Reference (PPER)	:	EN 420:2003+A1:2009	
\sim		EN ISO 374-1:2016+A1:2018	
		EN ISO 374-5:2016	

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 Sdn Bhd (75398-к)

No.7, Kawasan Perusahaan Suria 45600 Bestari Jaya Selangor Darul Ehsan, Malaysia Tel: +603 - 3280 3888 Fax: +603 - 3271 0135



We, Hartalega 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 Sdn. Bhd.

Place and Date of Issue

: Hartalega Sdn. Bhd./ 13th December 2021

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

Name : NURUL AISYAH KONG Position : DEPUTY GENERAL MANAGER – QUALITY ASSURANCE

ATTACHMENT I

Product or Trade Name	Reference Number
Peppler Sensitive Desire 207	XS: 207XS S: 207S M: 207M L: 207L XL:207XL

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		

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ATTACHMENT III

Product or Trade Name	Reference Number
Peppler Sensitive Desire 207	XS: 207XS S: 207S M: 207M L: 207L XL:207XL