

EU DECLARATION OF CONFORMITY

as per

PPE Regulation (EU) 2016/245 on Personal Protective Equipment

Manufacturer's Name : QUBE MEDICAL PRODUCTS SDN. BHD.
Manufacturer's Address : No. 9, Jalan KPK 1/3, Kawasan Perindustrian Kundang, 48020
Kundang Jaya, Selangor Darul Ehsan, Malaysia.
European Authorized : QUBE Integrated Europe GmbH
Representative : Lilienthalstr. 14,
41515, Grevenbroich
Germany

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

We hereby declare that the Personal Protective Equipment (PPE) described below is in conformity with the relevant Union harmonisation legislation, Regulation (EU) 2016/425 on Personal Protective Equipment, and complies with the applicable harmonised standards; EN 21420:2020+A1:2024, EN ISO 374-1:2016+A1:2018 & EN ISO 375-5:2016.

Product Name : Powder Free Latex Examination Gloves – Natural
Product Reference : NR9PFPTNPC
Description : Natural Rubber Latex Powder Free
Classification : Category III (Type C)
EU Type Examination : 2777/27019-02/E00-00
Certificate No.
Issued by : SATRA Technology Europe Ltd.
Bracetown Business Park,
Clonee,
Dublin D15 YN2P,
Ireland
Notified Body No. : 2777

The PPE is subject to the conformity assessment procedure based on quality assurance of the production process (Module D) under the surveillance of:

Module D Certificate No. :
Issued by : BSI Group The Netherlands B.V.
John M. Keynesplein 9,
1066 EP Amsterdam,
The Netherlands
Notified Body Number : 2797

(Place and date)

(Authorized signature & stamp)

Selangor, Malaysia on 25th May 2026


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Thong Seng Choon (Senior QA Executive)