



M A X T E R
GLOVE MANUFACTURING SDN BHD
(229862-H)

Lot 6070, Jalan Haji Abdul Manan
6th Miles Off Jalan Meru
41050 Klang, Selangor, Malaysia
Tel: 603-33929888 (8 lines) Fax: 603-33923328
E-MAIL: maxter@tm.net.my
www.maxter.com.my

Date: 19th May 2022

To Whom It May Concern:

EU DECLARATION OF CONFORMITY

We, **MAXTER GLOVE MANUFACTURING SDN. BHD.**, located at Lot 6070, Jalan Haji Abdul Manan, 6th Miles Off Jalan Meru, 41050 Klang, Selangor, Malaysia, declares under our sole responsibility that the medical devices and PPE described hereafter as:-

- **“PrimaDent” label, Latex Examination Gloves Powder Free**
Product Reference: **MTCLPF**
Basic UDI-DI: **955 500211 637CR**
Single Registration Number (SRN): **MY-MF-000016719**

are in conformity with:-

- The general safety and performance requirements of Annex I Medical Device Regulation (EU) 2017/745 for Class I medical devices.
- Classification: Class I based on Rule 5 transient use, Annex VIII of the Medical Device Regulation (EU) 2017/745
- With the national standard transposing harmonized standard EN455-1, EN455-2, EN455-3 and EN455-4 and is self-certified as a Class I non-sterile medical device.
- EN ISO 15223-1, EN ISO 20417, EN ISO 10993-1:2018, EN ISO 14971:2019 standards.
- The provisions of Personal Protective Equipment (PPE) – Regulation (EU) 2016/425 and the requirements of the European harmonized standard EN420:2003+A1:2009, EN ISO 374-1:2016, and EN ISO 374-5:2016.
- are PPE Category III covered by EU Type Examination (Module B)-certificate no.: **2777/12719-01/E00-00** issued by the Notified Body:
SATRA (2777)
Bracetown Business Park, Clonee D15YN2P, Republic of Ireland.
- Is subject to the conformity assessment procedure set out in Module D of Regulation (EU) 2016/425 as a Category III product and under the surveillance of the Notified Body:
SGS FIMKO OY (0598)
Takomotie 8, FI-00380 Helsinki, Finland.
- The gloves are manufactured according to ISO 9001:2015 and ISO 13485:2016 Quality Management Systems and certified by Notified Body, SGS United Kingdom Ltd Systems & Services Certification.
- Our Authorized EU Representative is Supermax Healthcare (Europe) Limited, 38 Main Street, Swords, Co. Dublin, Ireland K67 E0A2

Klang, Selangor
Malaysia


Yap Peak Geeh
QA & Regulatory Affairs Manager