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Date: 13th April 2023

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 described hereafter as:-

➤ "SUPERGLOVES" label, Non Sterile Powdered Latex Examination Gloves Basic UDI-DI: 955 500211 636CP

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
- The national standard transposing harmonized standard EN455-1, EN455-2, EN455-3 and EN455-4
- 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 European authorised representative is Supermax Healthcare (Europe) Limited, 38 Main Street, Swords, Co. Dublin, Ireland K67 E0A2

Signed for and on behalf of

Maxter Glove/Manufacturing Sdn Bhd

Yap Peak Geeh

QA & Regulatory Affairs Senior Manager

Klang, Selangor Malaysia