EU DECLARATION OF CONFORMITY

The Manufacturer ANSELL HEALTHCARE EUROPE N.V. RIVERSIDE BUSINESS PARK, BLOCK J BOULEVARD INTERNATIONAL 55 B-1070 BRUSSELS BELGIUM

declares under his sole responsibility, that the PPE described hereafter:

BioClean™ Nitramax BNMS Sterile Nitrile Cleanroom Gloves

Products manufactured as of: [2022/12/20]

PPE to be used against category III risks



is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN 421:2010, EN ISO 21420:2020, EN ISO 374-1:2016, EN ISO 374-5:2016 and is identical to the PPE which is subject to the EU-Type examination (Module B, Annex V of the Regulation), under certificate number 032/2022/1064, issued by the Notified Body:

CENTEXBEL (0493) TECHNOLOGIEPARK 70 B-9052 ZWIJNAARDE BELGIUM

and is subject to the procedure set out in Annex VIII (Module D) of the Regulation under the supervision of the Notified Body:

CENTEXBEL (0493) TECHNOLOGIEPARK 70 B-9052 ZWIJNAARDE BELGIUM

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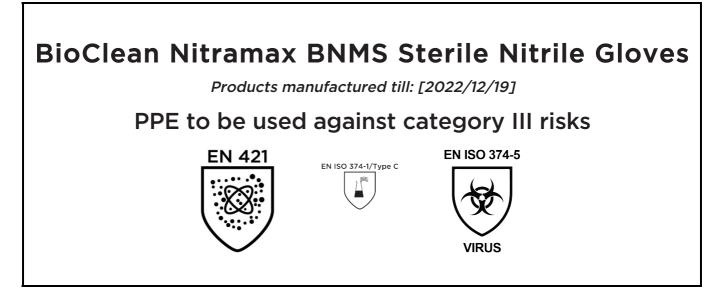
Guido Van Duren Director - Regulatory affairs Ansell

Place: Brussels Date: 2022/12/20

EU DECLARATION OF CONFORMITY

The Manufacturer NITRITEX (M) SDN BHD, NO.2, JALAN JURUNILAI U1/20, SEKSYEN U1, HICOM GLENMARIE INDUSTRIAL PARK, 40150 SHAH ALAM, SELANGOR, MALAYSIA

declares under his sole responsibility, that the PPE described hereafter:



is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN 420:2003 + A1:2009, EN 421:2010, EN ISO 374-1:2016, EN ISO 374-5:2016 and is identical to the PPE which is subject to the EU-Type examination (Module B, Annex V of the Regulation), under certificate number 060/2018/1754, issued by the Notified Body:

CENTEXBEL (0493) TECHNOLOGIEPARK 70 B-9052 ZWIJNAARDE BELGIUM

and is subject to the procedure set out in Annex VIII (Module D) of the Regulation under the supervision of the Notified Body:

SGS FIMKO OY (0598) TAKOMOTIE 8, FI-00380 HELSINKI, FINLAND

8 to Or

Guido Van Duren Director - Regulatory affairs Ansell

Place: Brussels Date: 2019/04/01