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 Excell BEXS Sterile Nitrile Gloves

Products manufactured as of: [2024/08/28]

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 032/2022/1083, 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

Guido Van Duren

Director - Regulatory affairs

Ansell

Place: Brussels Date: 2024/08/28

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 Excell BEXS Sterile Nitrile Cleanroom Gloves

Products manufactured as of: [2024/01/17] and till: [2024/08/27]

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 032/2022/1083, 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

Guido Van Duren

Director - Regulatory affairs

Ansell

Place: Brussels Date: 2024/01/17

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:

BioClean Excell BEXS Sterile Nitrile Gloves

Products manufactured till: [2024/01/16]

PPE to be used against category III risks

EN 421



EN ISO 374-5



EN ISO 374-1/Type C

is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN 421:2010, EN ISO 374-5:2016, EN ISO 374-1:2016, EN 420:2003 + A1:2009 and is identical to the PPE which is subject to the EU-Type examination (Module B, Annex V of the Regulation), under certificate number 060/2019/0070, 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**

Guido Van Duren

Director - Regulatory affairs

Ansell

Place: Brussels Date: 2019/04/01