欧盟符合性声明

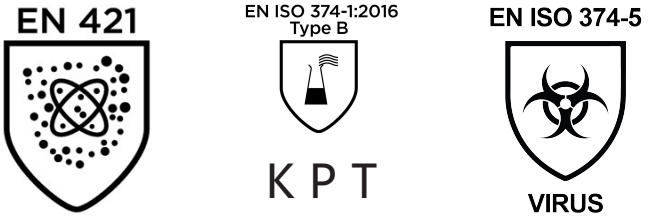
制造商
ANSELL HEALTHCARE EUROPE N.V.
RIVERSIDE BUSINESS PARK, BLOCK J
BOULEVARD INTERNATIONAL 55
B-1070 BRUSSELS
BELGIUM
WWW.ANSELL.COM

声明以下所述的个人防护设备由其全权负责：

BioClean™ Legacy BLA2 Non-sterile Latex Cleanroom Glove

适用产品起始日期 [2024/12/11]

用于防护category III风险的PPE



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+A1:2018, 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 032/2024/0713, issued by the Notified Body:

CENTEXBEL (0493)
TECHNOLOGIEPARK 70
B-9052 ZWIJNAARDE
BELGIUM

并须遵守该规例附件VI（模块D）所载的程序：

Guido Van Duren
Director - Regulatory affairs
Ansell

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B-9052 ZWIJNAARDE
BELGIUM

地点： Brussels
日期： 2024/12/11

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RIVERSIDE BUSINESS PARK, BLOCK J
BOULEVARD INTERNATIONAL 55
B-1070 BRUSSELS
BELGIUM
WWW.ANSELL.COM

声明以下所述的个人防护设备由其全权负责：

BioClean™ Legacy BLA2 Non-sterile Latex Cleanroom Glove

适用产品起始日期 [2024/07/16] and till: [2024/12/10]

用于防护category III风险的PPE

is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN 421:2010, EN ISO 21420:2020, EN ISO 374-5:2016, EN ISO 374-1:2016 and is identical to the PPE which is subject to the EU-Type examination (Module B, Annex V of the Regulation), under certificate number 032/2023/0011, issued by the Notified Body:

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Guido Van Duren
Director - Regulatory affairs
Ansell

地点： Brussels
日期： 2024/07/16

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制造商
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RIVERSIDE BUSINESS PARK, BLOCK J
BOULEVARD INTERNATIONAL 55
B-1070 BRUSSELS
BELGIUM
WWW.ANSELL.COM

声明以下所述的个人防护设备由其全权负责：

BioClean™ Legacy BLA2 Non-sterile Latex Cleanroom Glove

适用产品起始日期 [2023/01/06] and till: [2024/07/15]

用于防护category III风险的PPE

is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN 421:2010, EN ISO 21420:2020, EN ISO 374-5:2016, EN ISO 374-1:2016 and is identical to the PPE which is subject to the EU-Type examination (Module B, Annex V of the Regulation), under certificate number 032/2023/0011, issued by the Notified Body:

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TECHNOLOGIEPARK 70
B-9052 ZWIJNAARDE
BELGIUM

并须遵守该规例附件VI（模块D）所载的程序：

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

Guido Van Duren
Director - Regulatory affairs
Ansell

地点： Brussels
日期： 2023/01/06

欧盟符合性声明

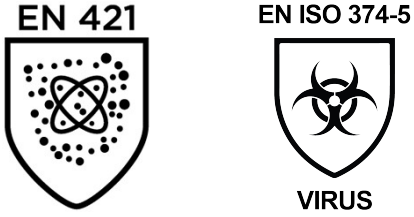
制造商
NITRITEX (M) SDN BHD,
NO.2, JALAN JURUNILAI U1/20,
SEKSYEN U1, HICOM GLENMARIE
INDUSTRIAL PARK,
40150 SHAH ALAM,
SELANGOR, MALAYSIA

声明以下所述的个人防护设备由其全权负责：

BioClean Legacy Non-sterile Latex Gloves BLA2

适用期至： [2023/01/05]

用于防护category III风险的PPE



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 374-5:2016 and is identical to the PPE which is subject to the EU-Type examination (Module B, Annex V of the Regulation), under certificate number 060/2018/1776, issued by the Notified Body:

CENTEXBEL (0493)
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B-9052 ZWIJNAARDE
BELGIUM

并须遵守该规例附件VI（模块D）所载的程序：

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

Guido Van Duren
Director - Regulatory affairs
Ansell

地点： Brussels
日期： 2019/04/01