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

声明以下所述的个人防护设备由其全权负责：

ActivArmr® (Class 00) RIG0011R

适用产品起始日期 [2024/08/12]

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 60903:2003, EN ISO 21420:2020 and is identical to the PPE which is subject to the EU-Type examination (Module B, Annex V of the Regulation), under certificate number 032/2023/0081.03, 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：

CENTEXBEL (0493)
TECHNOLOGIEPARK 70
B-9052 ZWIJNAARDE
BELGIUM

Guido Van Duren
Director - Regulatory affairs
Ansell

地点： Brussels
日期： 2024/08/11

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

声明以下所述的个人防护设备由其全权负责：

ActivArmr® (Class 00) RIG0011R

适用产品起始日期 [2024/06/06] and till: [2024/08/11]

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 60903:2003, EN ISO 21420:2020 and is identical to the PPE which is subject to the EU-Type examination (Module B, Annex V of the Regulation), under certificate number 032/2023/0081.02, issued by the Notified Body:

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Guido Van Duren
Director - Regulatory affairs
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地点： Brussels
日期： 2024/06/06

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BOULEVARD INTERNATIONAL 55
B-1070 BRUSSELS
BELGIUM

声明以下所述的个人防护设备由其全权负责：

ActivArmr® (Class 00) RIG0011R

适用期至： [2024/06/05]

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 60903:2003, EN ISO 21420:2020 and is identical to the PPE which is subject to the EU-Type examination (Module B, Annex V of the Regulation), under certificate number 032/2023/0081, issued by the Notified Body:

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Guido Van Duren
Director - Regulatory affairs
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地点： Brussels
日期： 2023/01/24