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

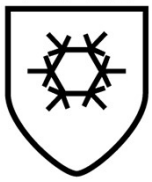
声明以下所述的个人防护设备由其全权负责：

ActivArmr® 97-631

适用产品起始日期 [2022/03/18]

用于防护category II风险的PPE

EN 511



12X

EN388: 2016

2231B

is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN 511:2006, EN 388:2016 +A1:2018, 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 032/2022/0426, issued by the Notified Body:

CENTEXBEL (0493)
TECHNOLOGIEPARK 70
B-9052 ZWIJNAARDE
BELGIUM

符合法规（欧盟）2016/425的规定和欧洲统一标准EN 511:2006, EN 388:2016 +A1:2018, EN ISO 21420:2020，并与进行EC型式检验的个人防护设备（PPE）相同；根据公告机构颁发的证书编号

Guido Van Duren
Director - Regulatory affairs
Ansell

地点： Brussels
日期： 2022/03/18

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B-1070 BRUSSELS
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声明以下所述的个人防护设备由其全权负责：

ActivArmr® 97-631

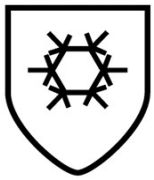
适用期至： [2022/03/17]

用于防护category II风险的PPE

EN 388

2231B

EN 511



12X

is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN 388:2016, EN 511:2006, 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 032/2018/0778, issued by the Notified Body:

CENTEXBEL (0493)
TECHNOLOGIEPARK 70
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符合法规（欧盟）2016/425的规定和欧洲统一标准EN 388:2016, EN 511:2006, EN 420:2003 + A1:2009，并与进行EC型式检验的个人防护设备（PPE）相同；根据公告机构颁发的证书编号

Guido Van Duren
Director - Regulatory affairs
Ansell

地点： Brussels
日期： 2018/04/25