

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：

HyFlex® 11-550

适用产品起始日期 [2021/11/16]

用于防护category III风险的PPE

EN388: 2016



4X21B

EN 407



X1XXXX

is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN 388:2016 +A1:2018, EN407:2020 , 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 032/2021/1196, 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

**CENTEXBEL (0493)
TECHNOLOGIEPARK 70
B-9052 ZWIJNAARDE
BELGIUM**

地点： Brussels
日期： 2021/11/16

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声明以下所述的个人防护设备由其全权负责：

HyFlex® 11-550

适用期至： [2021/11/15]

用于防护category III风险的PPE

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is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN 388:2016, EN 407:2004, 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 032/2018/2002.02, issued by the Notified Body:

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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：



BSI (0086)
KITEMARK COURT DAVY AVENUE KNOWLHILL
MILTON KEYNES MK5 8PP UNITED KINGDOM

Guido Van Duren
Director - Regulatory affairs
Ansell

地点： Brussels
日期： 2018/11/19