

## 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)  
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B-9052 ZWIJNAARDE  
BELGIUM

地点： Brussels  
日期： 2021/11/16

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**HyFlex® 11-550**

适用期至： [2021/11/15]

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4X21B



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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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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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地点： Brussels  
日期： 2018/11/19