

## 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：

**HyFlex® 11-754**

适用产品起始日期 [2022/01/28]

用于防护category II风险的PPE

EN388: 2016



**3X41D**

is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards 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 032/2022/0083, 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 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032/2022/0083：

Guido Van Duren  
Director - Regulatory affairs  
Ansell

地点： Brussels  
日期： 2022/01/28

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is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN 388:2016, 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 032/2020/1478, issued by the Notified Body:

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防护设备（PPE）相同；根据公告机构颁发的证书编号032/2020/1478：

Guido Van Duren  
Director - Regulatory affairs  
Ansell

地点： Brussels  
日期： 2020/12/04