

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® 11-591

适用产品起始日期 [2024/06/21]

用于防护category II风险的PPE

EN388: 2016



4X43F

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 032/2024/0139.02, 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, EN407:2020, EN ISO 21420:2020 并行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： 2024/06/21

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声明以下所述的个人防护设备由其全权负责：

HyFlex® 11-591

适用期至： [2024/06/20]

用于防护category III风险的PPE

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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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B-9052 ZWIJNAARDE
BELGIUM

地点： Brussels
日期： 2024/02/27