

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® 11-842

适用产品起始日期 [2024/06/14]

用于防护category II风险的PPE

EN388: 2016



4131A

EN 407



X1XXXX

is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN ISO 21420:2020, EN 388:2016 +A1:2018, EN407:2020 and is identical to the PPE which is subject to the EU-Type examination (Module B, Annex V of the Regulation), under certificate number 032/2022/0480.02, 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 ISO 21420:2020, EN 388:2016 +A1:2018, EN407:2020，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： 2024/06/14

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BELGIUM

声明以下所述的个人防护设备由其全权负责:

HyFlex® 11-842

适用期至: [2024/06/13]

用于防护category III风险的PPE

EN388: 2016



4131A

EN 407



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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

CENTEXBEL (0493)
TECHNOLOGIEPARK 70
B-9052 ZWIJNAARDE
BELGIUM

地点: Brussels
日期: 2022/03/25