# 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**

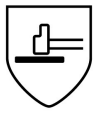
声明以下所述的个人防护设备由其全权负责：

## HyFlex® 11-812

*适用产品起始日期 [2022/01/28]*

用于防护category II风险的PPE

EN388: 2016



2010A

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 032/2022/0085, 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032/2022/0085：

Guido Van Duren
Director - Regulatory affairs
Ansell

地点： Brussels
日期： 2022/01/28

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Guido Van Duren
Director - Regulatory affairs
Ansell

地点： Brussels
日期： 2018/04/13