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：

ActivArmr® 97-012

适用产品起始日期 [2022/03/15]

用于防护category II风险的PPE

EN388: 2016

3131A

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 032/2022/0390, 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032/2022/0390：

Guido Van Duren
Director - Regulatory affairs
Ansell

地点： Brussels
日期： 2022/03/15

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：

ActivArmr® 97-012

适用产品起始日期 [2018/09/13] and till: [2022/03/14]

用于防护category II风险的PPE

EN 388

3131A

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 032/2018/1602, 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, EN 420:2003 + A1:2009，并与进入防护设备（PPE）相同；根据公告机构颁发的证书编号032/2018/1602：

Guido Van Duren
Director - Regulatory affairs
Ansell

地点： Brussels
日期： 2018/09/13

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：

ActivArmr® General Purpose Hi-Viz 97-012

适用产品起始日期 [2016/11/04] and till: [2018/09/12]

用于防护category II风险的PPE

EN 388

3131A

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 032/2016/1068, 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, EN 420:2003 + A1:2009，并与进入防护设备（PPE）相同；根据公告机构颁发的证书编号032/2016/1068：

Guido Van Duren
Director - Regulatory affairs
Ansell

地点： Brussels
日期： 2016/11/04

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：

ActivArmr® General Purpose Hi-Viz 97-012

适用期至： [2016/11/03]

用于防护category II风险的PPE

EN 388

3131

is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN 388:2003, EN 420:2003 + A1:2009 and is identical to the PPE which is subject to the EC Type examination; under certificate number 032/2014/0025 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 388:2003, EN 420:2003 + A1:2009，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032/2014/0025：

Guido Van Duren
Director - Regulatory affairs
Ansell

地点： Brussels
日期： 2014/01/20