

EU DECLARATION OF CONFORMITY

The Manufacturer

ANSELL HEALTHCARE EUROPE N.V.
RIVERSIDE BUSINESS PARK, BLOCK J
BOULEVARD INTERNATIONAL 55
B-1070 BRUSSELS
BELGIUM

declares under his sole responsibility, that the **PPE** described hereafter:

ActivArmr[®] 42-474

Products manufactured as of: [02/08/2018] and till: [02/08/2019]

PPE to be used against category III risks

EN 388:2016



2241B

EN 407



X2XXXX

is in conformity with the provisions of Regulation (EU) 2016/425 and with the European harmonized standards **EN 388:2016**, **EN 407:2004**, **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 032/2018/1335, issued by the Notified Body:

CENTEXBEL (0493)
TECHNOLOGIEPARK 70
B-9052 ZWIJNAARDE
BELGIUM

and is subject to the procedure set out in Annex VIII (Module **D**) of the Regulation under the supervision of the Notified Body:

BSI (0086)
KITEMARK COURT DAVY AVENUE KNOWLHILL
MILTON KEYNES MK5 8PP UNITED KINGDOM

Guido Van Duren
Director - Regulatory affairs
Ansell

Place: Brussels
Date: 02/08/2018

EU DECLARATION OF CONFORMITY

The Manufacturer

ANSELL HEALTHCARE EUROPE N.V.
RIVERSIDE BUSINESS PARK, BLOCK J
BOULEVARD INTERNATIONAL 55
B-1070 BRUSSELS
BELGIUM

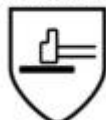
declares under his sole responsibility, that the **PPE** described hereafter:

ActivArmr® 42-474

Products manufactured as of: [03/08/2019]

PPE to be used against category III risks

EN 388:2016



2241B

EN 407



X2XXXX

is in conformity with the provisions of Regulation (EU) 2016/425 and with the European harmonized standards **EN 388:2016**, **EN 407:2004**, **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 032/2018/1335, issued by the Notified Body:

CENTEXBEL (0493)
TECHNOLOGIEPARK 70
B-9052 ZWIJNAARDE
BELGIUM

and is subject to the procedure set out in Annex VIII (Module D) of the Regulation under the supervision of the Notified Body:

CENTEXBEL (0493)
TECHNOLOGIEPARK 70
B-9052 ZWIJNAARDE
BELGIUM



Guido Van Duren
Director - Regulatory affairs
Ansell

Place: Brussels
Date: 03/08/2019

EU DECLARATION OF CONFORMITY

The Manufacturer

ANSELL HEALTHCARE EUROPE N.V.
RIVERSIDE BUSINESS PARK, BLOCK J
BOULEVARD INTERNATIONAL 55
B-1070 BRUSSELS
BELGIUM

declares under his sole responsibility, that the **PPE** described hereafter:

Crusader Flex 42-474

Products manufactured till: [13/12/2016]

PPE to be used against category III risks

EN 388:2003



2241

EN 407



X2XXXX

is in conformity with the provisions of Regulation (EU) 2016/425 and with the European harmonized standards **EN 388:2003**, **EN 407:2004**, **EN 420:2003 + A1:2009** and is identical to the **PPE** which is subject to the **EC** Type examination; under certificate number 03206267 issued by the Notified Body:

CENTEXBEL (0493)
TECHNOLOGIEPARK 70
B-9052 ZWIJNAARDE
BELGIUM

and is subject to the procedure set out in Annex VII (Module C2) of the Regulation under the supervision of the Notified Body:

CENTEXBEL (0493)
TECHNOLOGIEPARK 70
B-9052 ZWIJNAARDE
BELGIUM



Guido Van Duren
Director - Regulatory affairs
Ansell

Place: Brussels
Date: 02/06/2006