# 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® 46-511

*Products manufactured as of: [2018. 12. 06]*

PPE to be used against category III risks

EN 388



4X43DP

EN 407



X1XXXX

is in conformity with the provisions of Regulation (EU) 2016/425 and with the 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 032/2018/2094, 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: 2018. 12. 06

# EU DECLARATION OF CONFORMITY

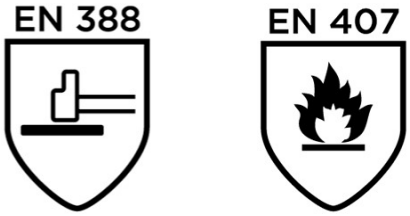
The Manufacturer
**ANSELL HEALTHCARE EUROPE N.V.**
**RIVERSIDE BUSINESS PARK, BLOCK J**
**BOULEVARD INTERNATIONAL 55**
**B-1070 BRUSSELS**
**BELGIUM**

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

## ActivArmr® 46-551

*Products manufactured till: [2018. 12. 05]*

PPE to be used against category III risks




4X43DP X1XXXX

is in conformity with the provisions of Regulation (EU) 2016/425 and with the 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 032/2016/1085, issued by the Notified Body:

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**BELGIUM**



Guido Van Duren
Director - Regulatory affairs
Ansell

Place: Brussels
Date: 2016. 11. 07