# 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

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**RIVERSIDE BUSINESS PARK, BLOCK J**
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**BELGIUM**

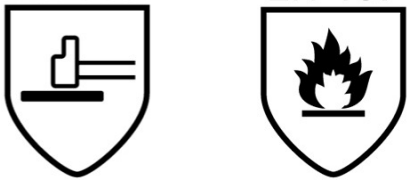
declares under his sole responsibility, that the PPE described hereafter:

## ActivArmr® 46-551

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

PPE to be used against category III risks

EN 388 EN 407



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Guido Van Duren
Director - Regulatory affairs
Ansell

Place: Brussels
Date: 2016/11/07