

# 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:

**ActivArm<sup>®</sup> 43-217**

**PPE to be used against category III risks**

EN388: 2016



**2111A**

EN 407



**41XX4X**

is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN 12477:2001 + A1:2005, EN 388:2016 +A1:2018, EN407:2020, 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 032/2023/0496, 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: 2023/08/08**

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**ANSELL HEALTHCARE EUROPE N.V.**  
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**B-1070 BRUSSELS**  
**BELGIUM**

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

**ActivArm<sup>®</sup> 43-217**

*Products manufactured as of: [2018/12/19]*

**PPE to be used against category III risks**

EN 407



41XX4X

EN 388



2121A

is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN 407:2004, EN 388:2016, EN 420:2003 + A1:2009, EN 12477:2001 + A1:2005 and is identical to the PPE which is subject to the EU-Type examination (Module B, Annex V of the Regulation), under certificate number 032/2018/2164, issued by the Notified Body:

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**B-9052 ZWIJNAARDE**  
**BELGIUM**

A handwritten signature in black ink, appearing to read 'Guido Van Duren'.

Guido Van Duren  
Director - Regulatory affairs  
Ansell

Place: Brussels  
Date: 2018/12/19

# EU DECLARATION OF CONFORMITY

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**ANSELL HEALTHCARE EUROPE N.V.**  
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**BELGIUM**

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

**ActivArmr<sup>®</sup> 43-217**

*Products manufactured till: [2018/12/18]*

**PPE to be used against category III risks**

EN 407



**41XX4X**

EN 388



**2121**

is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN 407:2004, EN 388:2003, EN 420:2003 + A1:2009, EN 12477:2001 + A1:2005 and is identical to the PPE which is subject to the EC Type examination; under certificate number 032/2016/0639 issued by the Notified Body:

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**Guido Van Duren**  
**Director - Regulatory affairs**  
**Ansell**

**Place: Brussels**  
**Date: 2016/07/29**