# EU DECLARATION OF CONFORMITY

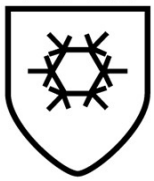
The Manufacturer
**ANSELL HEALTHCARE EUROPE N.V.**
**RIVERSIDE BUSINESS PARK, BLOCK J**
**BOULEVARD INTERNATIONAL 55**
**B-1070 BRUSSELS**
**BELGIUM**
**WWW.ANSELL.COM**

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

## ActivArmr™ 97-635

*Products manufactured as of: [2026/06/01]*

### PPE to be used against category II risks

| EN388: 2016 | EN 407 | EN 511 |
|---|---|---|
|  |  |  |
| 3X43E | X1XXXX | 021 |

is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN 388:2016 +A1:2018, EN407:2020 , EN 511:2006, 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 032/2025/0703.02, issued by the Notified Body:

**CENTEXBEL (0493)**
**TECHNOLOGIEPARK 70**
**B-9052 ZWIJNAARDE**
**BELGIUM**

and is subject to the procedure set out in Annex VI (Module C) of the Regulation.

Guido Van Duren
Director - Regulatory affairs
Ansell

Place: Brussels
Date: 2026/06/01

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## ActivArmr™ 97-635

*Products manufactured till: [2026/05/31]*

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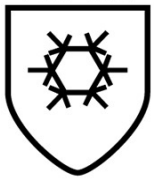
|  |  |  |
| 3X43E | X1XXXX | X11 |

is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN 388:2016 +A1:2018, EN407:2020 , EN 511:2006, 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 032/2025/0703, issued by the Notified Body:

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Guido Van Duren
Director - Regulatory affairs
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Place: Brussels
Date: 2025/11/21