

# 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® (Class 0) RIG011R**

*Products manufactured as of: [2024/08/12]*

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

is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN ISO 21420:2020, EN 60903:2003 and is identical to the PPE which is subject to the EU-Type examination (Module B, Annex V of the Regulation), under certificate number 032/2023/0081.03, 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: 2024/08/12**

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

## **ActivArmr® (Class 0) RIG011R**

*Products manufactured as of: [2024/06/06] and till: [2024/08/11]*

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Guido Van Duren  
Director - Regulatory affairs  
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Place: Brussels  
Date: 2024/06/06

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

**ActivArmr® (Class 0) RIG011R**

*Products manufactured till: [2024/06/05]*

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

**Place: Brussels**  
**Date: 2023/01/24**