

# 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<sup>®</sup> (Class 00) RIG0011R**

*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 60903:2003, 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 032/2023/0081.03, 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: 2024/08/11

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

**ActivArmr<sup>®</sup> (Class 00) RIG0011R**

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

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Date: 2024/06/06

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Guido Van Duren  
Director - Regulatory affairs  
Ansell

Place: Brussels  
Date: 2023/01/24