

# 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:

## AlphaTec™ 58-201

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

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

EN388: 2016



**3222B**

EN 407



**X2XXXX**

EN 511



**021**

EN ISO 374-1:2016  
Type A



**JKLOPT**

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, EN ISO 374-1:2016 and is identical to the PPE which is subject to the EU-Type examination (Module B, Annex V of the Regulation), under certificate number 032/2026/0287, 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: 2026/06/01

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## AlphaTec® 58-201

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

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

EN388: 2016



**3232B**

EN ISO 374-1:2016



**JKLOPT**

EN ISO 374-5:2016



**EN 407**



**X2XXXX**

**EN 511**



**021**

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Guido Van Duren  
Director – Regulatory affairs  
Ansell

Place: Brussels  
Date: 2022/08/12