

# EU DECLARATION OF CONFORMITY

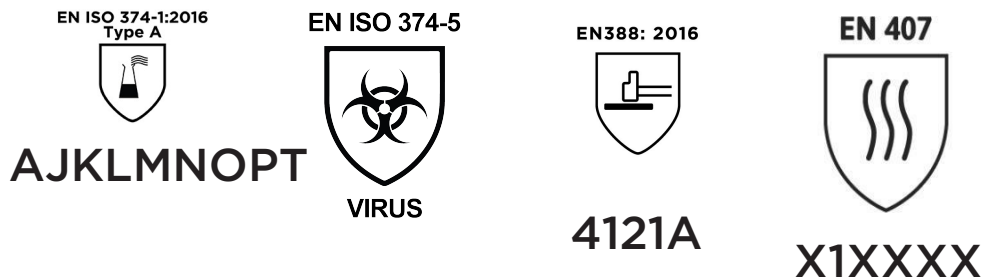
The Manufacturer  
**ANSELL HEALTHCARE EUROPE N.V.**  
**RIVERSIDE BUSINESS PARK, BLOCK J**  
**BOULEVARD INTERNATIONAL 55**  
**B-1070 BRUSSELS**  
**BELGIUM**  
**WWW.ANSELL.COM**

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

## **AlphaTec® 58-005**

*Products manufactured as of: [2025/07/29]*

### **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 ISO 374-1:2016, EN ISO 374-5:2016, EN 388:2016 +A1:2018, EN407:2020 and is identical to the PPE which is subject to the EU-Type examination (Module B, Annex V of the Regulation), under certificate number 032/2025/0474, 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: 2025/07/29**

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

## **AlphaTec® 58-005**

*Products manufactured as of: [2022/09/09] and till: [2025/07/28]*

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

EN388: 2016



**4122A**

EN 407



**X1XXXX**

EN ISO 374-1:2016  
Type A



**AJKLMNOPT**

EN ISO 374-5



**VIRUS**

is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN 388:2016 +A1:2018, EN407:2020 , EN ISO 21420:2020, EN ISO 374-1:2016, EN ISO 374-5:2016 and is identical to the PPE which is subject to the EU-Type examination (Module B, Annex V of the Regulation), under certificate number 032/2021/1090, issued by the Notified Body:

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**BELGIUM**



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

**Place: Brussels**  
**Date: 2022/09/09**

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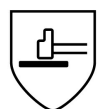
declares under his sole responsibility, that the PPE described hereafter:

## **AlphaTec® 58-005**

*Products manufactured as of: [2021/10/27] and till: [2022/09/08]*

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

EN388: 2016



**4122A**

EN 407



**X1XXXX**

EN ISO 374-1:2016  
Type A



**AJKLMNOPT**

EN ISO 374-5



**VIRUS**

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**BELGIUM**



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

**Place: Brussels**  
**Date: 2021/10/27**

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**ANSELL HEALTHCARE EUROPE N.V.**  
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BOULEVARD INTERNATIONAL 55  
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BELGIUM  
**WWW.ANSELL.COM**

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

## **AlphaTec® 58-005**

*Products manufactured till: [2021/10/26]*

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

**EN 388**



**4122A**

**EN 407**



**X1XXXX**

**EN ISO 374-1:2016  
Type A**



**AJKLMNOPT**

**EN ISO 374-5**



**VIRUS**

is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN 388:2016, EN 407:2004, EN 420:2003 + A1:2009, EN ISO 374-1:2016, EN ISO 374-5:2016 and is identical to the PPE which is subject to the EU-Type examination (Module B, Annex V of the Regulation), under certificate number 032/2020/1460, issued by the Notified Body:

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BELGIUM**

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

Place: Brussels  
Date: 2020/11/26