

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® 85-309

Products manufactured as of: [2025/02/07]

PPE to be used against category III risks

EN388: 2016



3101A

EN ISO 374-1:2016
Type B



AMP

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, 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 032/2021/1308.03, 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: 2025/02/07

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BOULEVARD INTERNATIONAL 55
B-1070 BRUSSELS
BELGIUM
WWW.ANSELL.COM

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

AlphaTec® 85-309

Products manufactured as of: [2023/12/20] and till: [2025/02/06]

PPE to be used against category III risks

EN388: 2016



3101A

EN ISO 374-1:2016
Type B



AMP

EN ISO 374-5



VIRUS

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CENTEXBEL (0493)
TECHNOLOGIEPARK 70
B-9052 ZWIJNAARDE
BELGIUM

and is subject to the procedure set out in Annex VII (Module C2) of the Regulation under the supervision of the Notified Body:

CENTEXBEL (0493)
TECHNOLOGIEPARK 70
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BELGIUM



Guido Van Duren
Director - Regulatory affairs
Ansell

Place: Brussels
Date: 2023/12/20

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B-1070 BRUSSELS
BELGIUM
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declares under his sole responsibility, that the PPE described hereafter:

AlphaTec® 85-309

Products manufactured as of: [2021/12/02] and till: [2023/12/19]

PPE to be used against category III risks

EN388: 2016



3101A

EN ISO 374-1:2016
Type B



AMP

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, 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 032/2021/1308, issued by the Notified Body:

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Guido Van Duren
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Place: Brussels
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declares under his sole responsibility, that the PPE described hereafter:

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Products manufactured till: [2021/12/01]

PPE to be used against category III risks

EN ISO 374-1:2016
Type B



AMP

EN ISO 374-5



VIRUS

EN 388



3101A

is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN ISO 374-1:2016 Type B, EN ISO 374-5:2016, EN 388:2016, EN 420:2003 + A1:2009 and is identical to the PPE which is subject to the EU-Type examination (Module B, Annex V of the Regulation), under certificate number 032/2019/0665.03, issued by the Notified Body:

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Guido Van Duren
Director - Regulatory affairs
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Place: Brussels
Date: 2021/01/12