

# 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:

**HyFlex™ 11-919**

*Products manufactured as of: [2026/04/01]*

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

EN388: 2016



**4121A**

EN 407



**X1XXXX**

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 and is identical to the PPE which is subject to the EU-Type examination (Module B, Annex V of the Regulation), under certificate number 032/2026/0175, issued by the Notified Body:

**CENTEXBEL (0493)**  
**TECHNOLOGIEPARK 70**  
**B-9052 ZWIJNAARDE**  
**BELGIUM**

and is subject to the procedure set out in Annex VI (Module C) of the Regulation.

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

**Place: Brussels**  
**Date: 2026/04/01**

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

**HyFlex® 11-919**

*Products manufactured as of: [2022/02/01] and till: [2026/03/31]*

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

EN388: 2016



**4121A**

is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN 388:2016 +A1:2018, 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 032/2022/0121, issued by the Notified Body:

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**TECHNOLOGIEPARK 70**  
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**BELGIUM**

and is subject to the procedure set out in Annex VI (Module C) of the Regulation.

A handwritten signature in black ink, appearing to read 'Guido Van Duren'.

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

**Place: Brussels**  
**Date: 2022/02/01**

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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:

**HyFlex® 11-919**

*Products manufactured as of: [2018/08/01] and till: [2022/01/31]*

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



is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards 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 032/2018/1324, issued by the Notified Body:

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A handwritten signature in black ink, appearing to read "Guido Van Duren".

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

**Place: Brussels**  
**Date: 2018/08/01**

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**BOULEVARD INTERNATIONAL 55**  
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**BELGIUM**  
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declares under his sole responsibility, that the PPE described hereafter:

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*Products manufactured till: [2018/07/31]*

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

EN 388



4121

is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN 388:2003, EN 420:2003 + A1:2009 and is identical to the PPE which is subject to the EC Type examination; under certificate number 032/2014/1275 issued by the Notified Body:

**CENTEXBEL (0493)**  
**TECHNOLOGIEPARK 70**  
**B-9052 ZWIJNAARDE**  
**BELGIUM**

and is subject to the procedure set out in Annex VI (Module C) of the Regulation.



Guido Van Duren  
Director - Regulatory affairs  
Ansell

Place: Brussels  
Date: 2014/09/18