# 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**

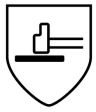
declares under his sole responsibility, that the PPE described hereafter:

## HyFlex® 11-725

*Products manufactured as of: [2022/03/23]*

PPE to be used against category II risks

EN388: 2016



4342B

is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN ISO 21420:2020, EN 388:2016 +A1:2018 and is identical to the PPE which is subject to the EU-Type examination (Module B, Annex V of the Regulation), under certificate number 032/2022/0457, 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: 2022/03/23

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

## HyFlex® 11-725

*Products manufactured till: [2022/03/22]*

PPE to be used against category II risks

EN 388

4342B

is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN 420:2003 + A1:2009, EN 388:2016 and is identical to the PPE which is subject to the EU-Type examination (Module B, Annex V of the Regulation), under certificate number 032/2019/1445, issued by the Notified Body:

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

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Guido Van Duren
Director - Regulatory affairs
Ansell

Place: Brussels
Date: 2019/09/23