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

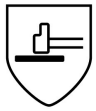
declares under his sole responsibility, that the PPE described hereafter:

HyFlex® 11-757

Products manufactured as of: [2022/02/04]

PPE to be used against category II risks

EN388: 2016



3X31F

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 032/2022/0142, 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: 2022/02/04

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

HyFlex® 11-757

Products manufactured till: [2022/02/03]

PPE to be used against category II risks

EN 388

3X31F

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 032/2021/0233, issued by the Notified Body:

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B-9052 ZWIJNAARDE
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Guido Van Duren
Director - Regulatory affairs
Ansell

Place: Brussels
Date: 2021/03/10