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® 11-571

Products manufactured as of: [2024/12/02]

PPE to be used against category II risks

EN 16350

EN388: 2016

4X43D

EN 407

X1XXXX

is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN 16350:2014, 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 032/2024/0697, 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: 2024/12/02

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Guido Van Duren
Director - Regulatory affairs
Ansell

Place: Brussels
Date: 2022/08/11