

EU DECLARATION OF CONFORMITY

The Manufacturer
ANSELL HEALTHCARE EUROPE N.V.
RIVERSIDE BUSINESS PARK, BLOCK J
BOULEVARD INTERNATIONAL 55
B-1070 BRUSSELS
BELGIUM

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

ActivArm[®] 97-631

Products manufactured as of: [2022/03/18]

PPE to be used against category II risks



12X

EN388: 2016



2231B

is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN 511:2006, EN 388:2016, 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 032/2022/0426, 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: 2022/03/18

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

ActivArm[®] 97-631

Products manufactured till: [2022/03/17]

PPE to be used against category II risks



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is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN 388:2016, EN 511:2006, 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 032/2018/0778, issued by the Notified Body:

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TECHNOLOGIEPARK 70
B-9052 ZWIJNAARDE
BELGIUM

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Guido Van Duren
Director - Regulatory affairs
Ansell

Place: Brussels
Date: 2018/04/25