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:

ActivArmr® 43-217

Products manufactured as of: [2018/12/19]

PPE to be used against category III risks

EN 407

41XX4X

EN 388

2121A

is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN 407:2004, EN 388:2016, EN 420:2003 + A1:2009, EN 12477:2001 + A1:2005 and is identical to the PPE which is subject to the EU-Type examination (Module B, Annex V of the Regulation), under certificate number 032/2018/2164, issued by the Notified Body:

CENTEXBEL (0493)
TECHNOLOGIEPARK 70
B-9052 ZWIJNAARDE
BELGIUM

and is subject to the procedure set out in Annex VII (Module C2) of the Regulation under the supervision of the Notified Body:

CENTEXBEL (0493)
TECHNOLOGIEPARK 70
B-9052 ZWIJNAARDE
BELGIUM

Guido Van Duren
Director - Regulatory affairs
Ansell

Place: Brussels
Date: 2018/12/19

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RIVERSIDE BUSINESS PARK, BLOCK J
BOULEVARD INTERNATIONAL 55
B-1070 BRUSSELS
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declares under his sole responsibility, that the PPE described hereafter:

ActivArmr® 43-217

Products manufactured till: [2018/12/18]

PPE to be used against category III risks

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Guido Van Duren
Director - Regulatory affairs
Ansell

Place: Brussels
Date: 2016/07/29