EU DECLARATION OF CONFORMITY

The manufacturer:

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

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

Powerflex® 80-400

applicable until [26-02-2019]

PPE to be used against category III risks

EN388:2003	EN407	EN511
X231	X2XXXX	020

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is in conformity with the provisions of Regulation (EU) 2016/425 and with the European harmonised standards EN420:2003+A1:2009, EN388:2003, EN407, EN511, and is identical to the PPE which is subject to the EC Type Examination; under certificate number 3206490 issued by the Notified Body:

CENTEXBEL (0493)
TECHNOLOGIEPARK 7
B-9052 ZWIJNAARDE

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 7
B-9052 ZWIJNAARDE

Guido Van Duren
Director – Regulatory Affairs PPE Products
Ansell

Date: 20-10-2006
Place: Brussels

EU DECLARATION OF CONFORMITY

The manufacturer:

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

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

ActivArmr 80-400

applicable as of [27-02-2019]

PPE to be used against category III risks

EN388:2016	EN407	EN511
2231C	X2XXXX	020

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is in conformity with the provisions of Regulation (EU) 2016/425 and with the European harmonised standards EN420:2003+A1:2009, EN388:2016, EN407, EN511, and is identical to the PPE which is subject to the EU Type Examination (Module B, Annex V of the Regulation); under certificate number 032/2019/0340 issued by the Notified Body:

CENTEXBEL (0493)
TECHNOLOGIEPARK 7
B-9052 ZWIJNAARDE

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

CENTEXBEL (0493)
TECHNOLOGIEPARK 7
B-9052 ZWIJNAARDE

Guido Van Duren
Director – Regulatory Affairs PPE Products
Ansell

Date: 27-02-2019
Place: Brussels