

Ansell Healthcare Europe N.V.

Riverside Business Park
Boulevard International 55

Block J
B-1070 Brussels

Tel. 32 (0)2-528 74 00
Fax 32 (0)2-528 74 01

EC DECLARATION OF PRODUCT CONFORMITY

Category III

The manufacturer, established in the European Economic Community:

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

declares that the PPE described hereafter:

ActivArmr® 80-813



EN 388: 2003
2542



412110

is in conformity with the provisions of the Council Directive 89/686/EEC and with the European harmonised standards EN420:2003+A1:2009, EN388: 2003 & EN407: 2004, and is identical to the PPE which is subject to the EC Type Examination certificate number 03211731 issued by the Notified Body:

**CENTEXBEL (0493)
TECHNOLOGIEPARK 7
B-9052 ZWIJNAARDE**

is subject to the procedure set out in Article 11 point A of Directive 89/686/EEC under the supervision of the Notified Body

**CENTEXBEL (0493)
TECHNOLOGIEPARK 7
B-9052 ZWIJNAARDE**

**Tuesday, 27 March 2018
Guido Van Duren
Director – Global Regulatory Affairs
PPE Products
Ansell**

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EC DECLARATION OF PRODUCT CONFORMITY

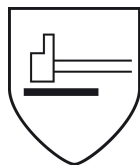
Category III

The manufacturer, established in the European Economic Community:

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

declares that the PPE described hereafter:

ActivArmr® 80-813



EN 388: 2016
2X42C



412110

is in conformity with the provisions of the Council Directive 89/686/EEC and with the European harmonised standards EN420:2003+A1:2009, EN388: 2016 & EN407: 2004, and is identical to the PPE which is subject to the EC Type Examination certificate number 032/2016/1096 issued by the Notified Body:

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B-9052 ZWIJNAARDE**

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