

Ansell Healthcare Europe N.V.

Riverside Business Park
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Block J
B-1070 Brussels

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EC DECLARATION OF PRODUCT CONFORMITY

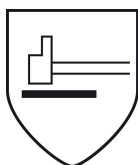
Category III

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® 43-217



EN 388: 2003
2121



41XX4X

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, EN12477:2001 + A1:2005, and is identical to the PPE which is subject to the EC Type Examination certificate number 032/2016/0639 issued by the Notified Body:

**CENTXBEL (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

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