

# 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:

## **VersaTouch® 78-103**

*applicable until [29-10-2018]*

**PPE to be used against category II risks**

EN388:2003



113X

EN511



010

is in conformity with the provisions of Regulation (EU) 2016/425 and with the European harmonised standards EN420:2003+A1:2009, EN388:2003, EN511, and is identical to the PPE which is subject to the EC Type Examination; under certificate number 03213603 issued by the Notified Body:

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



Guido Van Duren  
Director – Regulatory Affairs PPE Products  
Ansell

Date: 29-08-2013  
Place: Brussels

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**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® 78-103**

*applicable as of [30-10-2018]*

**PPE to be used against category II risks**

EN388:2016



113XB

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010

is in conformity with the provisions of Regulation (EU) 2016/425 and with the European harmonised standards EN420:2003+A1:2009, EN388:2016, 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 032/2018/1889 issued by the Notified Body:

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

and is subject to the procedure set out in Annex VI (Module C) of the Regulation.



Guido Van Duren  
Director – Regulatory Affairs PPE Products  
Ansell

Date: 30-10-2018  
Place: Brussels