# EU DECLARATION OF CONFORMITY

The manufacturer:

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

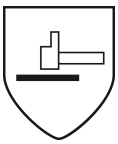
declares under his sole responsibility, that the PPE described hereafter:

## Powerflex® 80-100

*applicable until [11-04-2019]*

PPE to be used against category II risks

EN388:2003



2242

is in conformity with the provisions of Regulation (EU) 2016/425 and with the European harmonised standards EN420:2003+A1:2009, EN388:2003, and is identical to the PPE which is subject to the EC Type Examination; under certificate number 3210039 issued by the Notified Body:

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

Guido Van Duren
Director – Regulatory Affairs PPE Products
Ansell

Date: 29-01-2010
Place: Brussels

# EU DECLARATION OF CONFORMITY

The manufacturer:

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

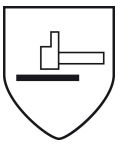
declares under his sole responsibility, that the PPE described hereafter:

## ActivArmr® 80-100

*applicable as of [12-04-2019]*

PPE to be used against category II risks

EN388:2016



2242B

is in conformity with the provisions of Regulation (EU) 2016/425 and with the European harmonised standards EN420:2003+A1:2009, EN388:2016, and is identical to the PPE which is subject to the EU Type Examination (Module B, Annex V of the Regulation); under certificate number 032/2019/0687 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: 12-04-2019
Place: Brussels