EU DECLARATION OF CONFORMITY

The manufacturer:

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

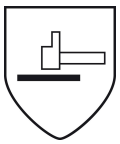
declares under his sole responsibility, that the PPE described hereafter:

Winter Hi Viz™ 23-491

applicable until [06-11-2018]

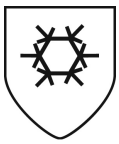
PPE to be used against category II risks

EN388:2003



3121

EN511



111

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 3205076 issued by the Notified Body:

CENTEXBEL (0493)
TECHNOLOGIEPARK 7
B-9052 ZWIJNAARDE

Guido Van Duren
Director – Regulatory Affairs PPE Products
Ansell

Date: 17-03-2005
Place: Brussels

EU DECLARATION OF CONFORMITY

The manufacturer:

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

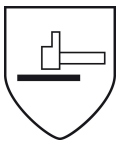
declares under his sole responsibility, that the PPE described hereafter:

ActivArmr® 23-491

applicable as of [07-11-2018]

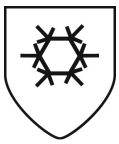
PPE to be used against category II risks

EN388:2016



3121B

EN511



111

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 032/2018/1928 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: 07-11-2018
Place: Brussels