EU DECLARATION OF CONFORMITY

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

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

ActivArmr[®] 97–003

Products manufactured as of: [2019/01/25]

PPE to be used against category II risks

is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN 420:2003 + A1:2009 and is identical to the PPE which is subject to the EU-Type examination (Module B, Annex V of the Regulation), under certificate number 032/2019/0140, issued by the Notified Body:



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

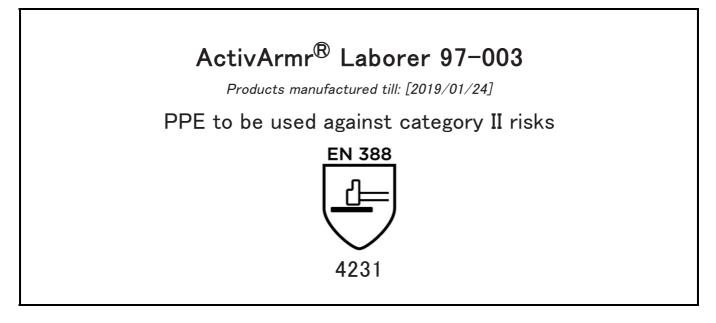
Guido Van Duren Director - Regulatory affairs Ansell

Place: Brussels Date: 2019/01/25

EU DECLARATION OF CONFORMITY

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

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



is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN 388:2003, EN 420:2003 + A1:2009 and is identical to the PPE which is subject to the EC Type examination; under certificate number 032/2017/0153 issued by the Notified Body:

CENTEXBEL (0493) TECHNOLOGIEPARK 70 B-9052 ZWIJNAARDE BELGIUM

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

Guido Van Duren Director - Regulatory affairs Ansell

Place: Brussels Date: 2017/02/06