EU DECLARATION OF CONFORMITY

The manufacturer:

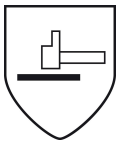
ANSELL HEALTHCARE EUROPE N.V.
RIVERSIDE BUSINESS PARK, BLOCK J
BOULEVARD INTERNATIONAL 55
B-1070 BRUSSELS

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

HyFlex® 11-542

PPE to be used against category III risks

EN388:2016



4X32F

EN407

X1XXXX

is in conformity with the provisions of Regulation (EU) 2016/425 and with the European harmonised standards EN420:2003+A1:2009, EN388:2016, EN407, and is identical to the PPE which is subject to the EU Type Examination (Module B, Annex V of the Regulation); under certificate number 032/2018/1414 issued by the Notified Body:

CENTEXBEL (0493)
TECHNOLOGIEPARK 7
B-9052 ZWIJNAARDE

and is subject to the procedure set out in Annex VII (Module D) of the Regulation under the supervision of the Notified Body:

BSI (0086)
Kitemark Court Davy Avenue Knowlhill
Milton Keynes MK5 8PP United Kingdom

Guido Van Duren
Director – Regulatory Affairs PPE Products
Ansell

Date: 16-08-2018
Place: Brussels