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:

HyFlex® 11-130

Products manufactured as of: [2018/04/23]

PPE to be used against category II risks

EN 388

3131A

EN 16350

is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN 388:2016, EN 420:2003 + A1:2009, EN 16350:2014 and is identical to the PPE which is subject to the EU-Type examination (Module B, Annex V of the Regulation), under certificate number 032/2018/0747, 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: 2018/04/23

EU DECLARATION OF CONFORMITY

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

and authorized representative:
COMASEC S.A.S
5 ALLÉE DES BAS TILLIERS
92238 GENNEVILLIERS CEDEX
FRANCE

declare under their sole responsibility, that the PPE described hereafter:

PU 610 AS

Products manufactured till: [2018/04/22]

PPE to be used against category II risks

EN 388

3121

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 0072/015/162/02/07/0010 issued by the Notified Body:

IFTH – INSTITUT FRANÇAIS TEXTILE-HABILLEMENT (0072)
AVENUE GUY DE COLLONGUE – 69134 ECULLY CEDEX – FRANCE

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: 2007/02/21