## **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**<sup>®</sup> 11-800

Products manufactured till: [07/12/2016]

PPE to be used against category II risks



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is in conformity with the provisions of Regulation (EU) 2016/425 and with the European harmonized standards EN 388:2003, EN 420:2003 + A1:2009, EN 420:2010 and is identical to the PPE which is subject to the EC Type examination; under certificate number 03205160 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: 02/05/2005

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ANSELL HEALTHCARE EUROPE N.V.
RIVERSIDE BUSINESS PARK, BLOCK J
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declares under his sole responsibility, that the PPE described hereafter:

## **HyFlex**<sup>®</sup> 11-800

Products manufactured as of: [08/12/2016]

PPE to be used against category II risks

EN 388:2016



is in conformity with the provisions of Regulation (EU) 2016/425 and with the European harmonized standards EN 388:2016, EN 420:2003 + A1:2009 and is identical to the PPE which is subject to the EC-Type examination (Module B, Annex V of the Regulation), under certificate number 032/2016/1244, 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: 08/12/2016