## **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® 11-840**

PPE to be used against category II risks

EN388:2003



is in conformity with the provisions of Regulation (EU) 2016/425 and with the European harmonised standards EN420:2003+A1:2009, EN388:2003, and is identical to the PPE which is subject to the EC Type Examination; under certificate number 03213141 issued by the Notified Body:

> **CENTEXBEL (0493) TECHNOLOGÌEPAŔK 7 B-9052 ZWIJNAARDE**

Guido Van Duren

Director - Regulatory Affairs PPE Products

Ansell

Date: 29-01-2019 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:

## **HyFlex® 11-840**

PPE to be used against category II risks

EN388:2016



4131A

is in conformity with the provisions of Regulation (EU) 2016/425 and with the European harmonised standards EN420:2003+A1:2009, EN388:2016, and is identical to the PPE which is subject to the EU Type Examination (Module B, Annex V of the Regulation); under certificate number 032/2017/1106 issued by the Notified Body:

> **CENTEXBEL (0493) TECHNOLOGÌEPAŔK 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: 29-01-2019 Place: Brussels