

# 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:

## **Microflex® 93-850**

**PPE to be used against category III risks**

EN ISO 374-1:2016

Type B



JKOPT

EN ISO 374-5:2016



VIRUS

EN421:2010



is in conformity with the provisions of Regulation (EU) 2016/425 and with the European harmonised standards EN420:2003+A1:2009, EN ISO 374:2016, EN421:2010, and is identical to the PPE which is subject to the EU Type Examination (Module B, Annex V of the Regulation); under certificate number 032/2018/1558 issued by the Notified Body:

**CENTEXBEL (0493)  
TECHNOLOGIEPARK 7  
B-9052 ZWIJNAARDE**

and is subject to the procedure set out in Annex VII (Module C2) of the Regulation under the supervision of the Notified Body:

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Guido Van Duren  
Director – Regulatory Affairs PPE Products  
Ansell

Date: 18-09-2018  
Place: Brussels