

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:
MicroFlex® 93-843

Products manufactured as of: [21/04/2018]

PPE to be used against category III risks

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is in conformity with the provisions of Regulation (EU) 2016/425 and with the European harmonized standards EN 420:2003 + A1:2009, EN ISO 374-1:2016, EN ISO 374-5:2016 and is identical to the PPE which is subject to the EU-Type examination (Module B, Annex V of the Regulation), under certificate number 032/2018/0291, issued by the Notified Body:

CENTEXBEL (0493)
TECHNOLOGIEPARK 70
B-9052 ZWIJNAARDE
BELGIUM

and is subject to the procedure set out in Annex VII (Module C2) of the Regulation under the supervision of the Notified Body:
CENTEXBEL (0493)

TECHNOLOGIEPARK 70
B-9052 ZWIJNAARDE
BELGIUM

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Guido Van Duren
Director - Regulatory affairs
Ansell
Place: Brussels
Date: 14/02/2018

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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:
MicroFlex® 93-843

Products manufactured till: [20/04/2018]

PPE to be used against category III risks

EN374_LowChemicalResistance

EN 374 Microorganism

is in conformity with the provisions of Regulation (EU) 2016/425 and with the European harmonized standards EN 374:2003 , EN 420:2003 + A1:2009 and is identical to the PPE which is subject to the EC Type examination; under certificate number 032/2014/1449 issued by the Notified Body:
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Guido Van Duren

Director - Regulatory affairs

Ansell

Place: Brussels

Date: 21/10/2014