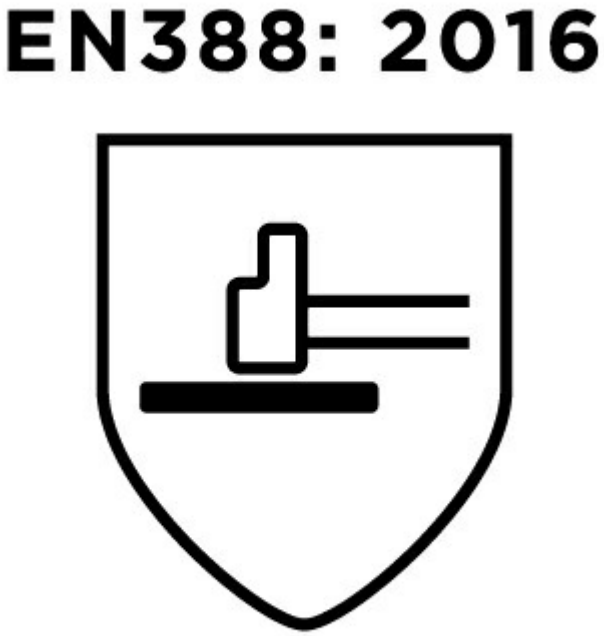
EU DECLARATION OF CONFORMITY

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® 11-423
Products manufactured as of: [2021/11/19]
PPE to be used against category III risks



4X32B

X1XXXX
is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN 388:2016 +A1:2018, EN407:2020 , EN ISO 21420:2020 and is identical to the PPE which is subject to the EU-Type examination (Module B, Annex V of the Regulation), under certificate number 032/2021/1224, issued by the Notified Body:
CENTEXBEL (0493)
TECHNOLOGIEPARK 70

B-9052 ZWIJNAARDE
BELGIUM
and is subject to the procedure set out in Annex VIII (Module D) of the Regulation under the supervision of the Notified Body:
CENTEXBEL (0493)
TECHNOLOGIEPARK 70
B-9052 ZWIJNAARDE
BELGIUM

Guido Van Duren
Director - Regulatory affairs
Ansell
Place: Brussels
Date: 2021/11/19

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® 11-423
Products manufactured as of: [2018/12/06] and till: [2021/11/18]
PPE to be used against category III risks

4X32B

X1XXXX
is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN 388:2016, EN 407:2004, EN 420:2003 + A1:2009 and is identical to the PPE which is subject to the EU-Type examination (Module B, Annex V of the Regulation), under certificate number 032/2018/2095, 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

Guido Van Duren
Director - Regulatory affairs
Ansell
Place: Brussels
Date: 2018/12/06

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® 11-423
Products manufactured till: [2018/12/05]
PPE to be used against category III risks

4332

X1XXXX
is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN 388:2003, EN 407:2004, EN 420:2003 + A1:2009 and is identical to the PPE which is subject to the EC Type examination; under certificate number 032/2014/1273 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

Guido Van Duren
Director - Regulatory affairs
Ansell
Place: Brussels
Date: 2014/09/18