# 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:

## AccuTech® 91-210

*Products manufactured till: [2026/12/16]*

### PPE to be used against category III risks

EN ISO 374-5

VIRUS

EN ISO 374-1:2016
Type B

KPT

EN 421

is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN ISO 374-5:2016, EN ISO 21420:2020, EN ISO 374-1:2016, EN 421:2010 and is identical to the PPE which is subject to the EU-Type examination (Module B, Annex V of the Regulation), under certificate number 032/2021/1392, 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: 2021/12/16

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declares under his sole responsibility, that the PPE described hereafter:

## AccuTech® 91-210

### PPE to be used against category III risks

is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN 420:2003 + A1:2009, EN ISO 374-1:2016 Type B, EN ISO 374-5:2016, EN 421:2010 and is identical to the PPE which is subject to the EU-Type examination (Module B, Annex V of the Regulation), under certificate number 032/2020/1042, 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: 2020/08/04