

# UK 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:

## **BioClean™ Legacy BLA2 Non-sterile Latex Cleanroom Glove**

*Products manufactured as of: [2024/12/11]*

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



**K P T**

**VIRUS**

is in conformity with the provisions of Regulation 2016/425 on personal protective equipment, as amended to apply in GB, and with the standards EN 421:2010, EN ISO 21420:2020, EN ISO 374-1:2016+A1:2018, EN ISO 374-5:2016 and is identical to the PPE which is subject to the Type-examination (Module B, Annex V of the Regulation), under certificate number 032/2024/0713, issued by the Body:

**CENTEXBEL (0493)**  
**TECHNOLOGIEPARK 70**  
**B-9052 ZWIJNAARDE**  
**BELGIUM**

and is subject to the conformity assessment procedure set in out in Annex VIII (Module D) of the Regulation under the surveillance of the Body:

**CENTEXBEL (0493)**  
**TECHNOLOGIEPARK 70**  
**B-9052 ZWIJNAARDE**  
**BELGIUM**

Guido Van Duren  
Director - Regulatory affairs  
Ansell

Place: Brussels  
Date: 2024/12/11

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

## **BioClean™ Legacy BLA2 Non-sterile Latex Cleanroom Glove**

*Applicable Until [2024/12/10]*

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

**EN 421**



EN ISO 374-1/Type C



**EN ISO 374-5**



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Guido Van Duren  
Director - Regulatory affairs  
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Place: Brussels  
Date: 2023/01/06