

# UK DECLARATION OF CONFORMITY

The Manufacturer  
**ANSELL HEALTHCARE EUROPE N.V.**  
**RIVERSIDE BUSINESS PARK, BLOCK J**  
**BOULEVARD INTERNATIONAL 55**  
**B-1070 BRUSSELS**  
**BELGIUM**

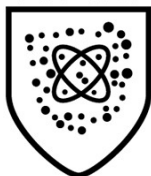
declares under his sole responsibility, that the PPE described hereafter:

## **BioClean™ Legion BLA3 Non-sterile Latex Cleanroom Glove**

*Applicable Until [2028/06/06]*

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

**EN 421**



**EN ISO 374-5**



**VIRUS**

**EN ISO 374-1/Type C**



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 374-5:2016, EN ISO 374-1:2016 and is identical to the PPE which is subject to the Type-examination (Module B, Annex V of the Regulation), under certificate number 032/2023/0012, 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 INTERNATIONAL LTD,**  
**8 NORTHUMBERLAND AVENUE,**  
**LONDON, WC2N 5BY,**  
**UK**

**Guido Van Duren**  
**Director - Regulatory affairs**  
**Ansell**

**Place: Brussels**  
**Date: 2023/01/06**