

# 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™ Sterile Nitrile RABS/Isolator Mitten** **GGL30NITM9**

*Applicable Until [2028/07/25]*

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

EN388: 2016



**4101X**

**EN 421**



EN ISO 374-1:2016  
Type A



**AJKNOST**

EN ISO 374-5



**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 388:2016 +A1:2018, EN 421:2010, EN ISO 21420:2020, EN ISO 374-1:2016, 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 032/2023/0463, 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:

Guido Van Duren  
Director - Regulatory affairs  
Ansell

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**8 NORTHUMBERLAND AVENUE,**  
**LONDON, WC2N 5BY,**  
**UK**

Place: Brussels  
Date: 2023/07/25