

# UK DECLARATION OF CONFORMITY

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

UK Importer  
**PATIENT GUARD LTD**  
**LANCASTER HOUSE, AMY JOHNSON**  
**WAY,**  
**BLACKPOOL, LANCASHIRE,**  
**FY4 2RP, UNITED KINGDOM**  
**INFO@PATIENTGUARD.CO.UK**

declara bajo su única responsabilidad que el EPI descrito a continuación:

**Edge® 48-919**

*Products manufactured as of: [2024/04/10]*

**PPE to be used against categoría II risks**

EN388: 2016



**4121A**

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 ISO 21420:2020 and is identical to the PPE which is subject to the UK Type-examination (Module B, Annex V of the Regulation), under certificate number AB0321/20670-02/E00-00, issued by the Approved Body:

**SATRA TECHNOLOGY CENTRE (0321)**  
**WYNDHAM WAY, TELFORD WAY,**  
**KETTERING, NORTHAMPTONSHIRE,**  
**NN16 8SD, UNITED KINGDOM**

y está sujeto al procedimiento establecido en el anexo VI (módulo C) del Reglamento.

A handwritten signature in black ink, appearing to read 'Guido Van Duren', written over a horizontal line.

Guido Van Duren  
Director - Regulatory affairs  
Ansell

Lugar: Bruselas  
Fecha: 2024/04/10

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*Applicable Until [2024/04/09]*

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Guido Van Duren  
Director - Regulatory affairs  
Ansell

Lugar: Bruselas  
Fecha: 2022/03/15