

# UK DECLARATION OF CONFORMITY

The Manufacturer  
**ANSELL HEALTHCARE EUROPE N.V.**  
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UK Importer  
**PATIENT GUARD LTD**  
LANCASTER HOUSE,  
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declare under their sole responsibility, that the PPE described hereafter:

**AphaTec® 58-530B**

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

EN388: 2016



**3122A**

EN 407



**X1XXXX**

EN ISO 374-1:2016  
Type A



**AJKLOPT**

EN ISO 374-5:2016



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, EN407:2020 , 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 UK Type-examination (Module B, Annex V of the Regulation), under certificate number AB0321/2256-01/E00-00, issued by the Approved Body:

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

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

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

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

Place: Brussels  
Date: 2022/10/11