

# 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:

**ActivArmr<sup>®</sup> 80-658**

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

EN388: 2016



**XX43E**

EN 407



**X2XXXX**

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 and is identical to the PPE which is subject to the UK Type-examination (Module B, Annex V of the Regulation), under certificate number AB0321/22871-01/E00-00, issued by the Approved Body:

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

and is subject to the conformity assessment procedure set out in Annex VII (Module C2) of the Regulation under the surveillance of the Approved Body:

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

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

**Place: Brussels  
Date: 2022.11.21**