# UK DECLARATION OF CONFORMITY

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

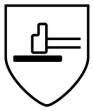
UK Importer
**PATIENT GUARD LTD**
**LANCASTER HOUSE,**
**AMY JOHNSON WAY,**
**BLACKPOOL, LANCASHIRE,**
**FY4 2RP, UNITED KINGDOM**
**INFO@PATIENTGUARD.CO.UK**

declare under their sole responsibility, that the PPE described hereafter:

## ActivArmr® 97-681

PPE to be used against category II risks

EN388: 2016



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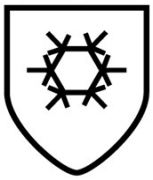
EN 511



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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 511:2006, 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 AB0321/24884-01/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 out in Annex VI (Module C) of the Regulation.

Guido Van Duren
Director – Regulatory affairs
Ansell

Place: Brussels
Date: 2023/09/28