

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:

MICROFLEX® 94-242

Products manufactured as of: [2024/02/07]

PPE to be used against category III risks

EN 16350



**EN ISO 374-1:2016
Type B**



J K P T

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 16350:2014, EN ISO 21420:2020, EN ISO 374-1:2016+A1:2018, 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 AB0321/24614-01/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 in out in Annex VIII (Module D) of the Regulation under the surveillance of the Body:

**CENTEXBEL (0493)
TECHNOLOGIEPARK 70
B-9052 ZWIJNAARDE
BELGIUM**

Guido Van Duren
Director - Regulatory affairs
Ansell

Place: Brussels
Date: 2024/02/07

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declares under his sole responsibility, that the PPE described hereafter:

MICROFLEX[®] 94-242

Applicable Until [2024/02/06]

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Type B



J K P T

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Guido Van Duren
Director - Regulatory affairs
Ansell

Place: Brussels
Date: 2023/08/10