

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:
TouchNTuff® 92-600
Products manufactured as of: [2024/12/06]
PPE to be used against category III risks

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J K P T

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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 ISO 21420:2020, EN ISO 374-1:2016+A1:2018, EN ISO 374-5:2016, ISO 18889:2019 and is identical to the PPE which is subject to the Type-examination (Module B, Annex V of the Regulation), under certificate number 032/2024/0500, issued by the Body:

CENTEXBEL (0493)
TECHNOLOGIEPARK 70
B-9052 ZWIJNAARDE
BELGIUM

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: 2024/12/06

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

TouchNTuff® 92-600

Applicable Until [2024/12/05]

PPE to be used against category III risks

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Guido Van Duren
Director - Regulatory affairs
Ansell
Place: Brussels
Date: 2024/04/15