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® 47-401

Products manufactured till: [2027/03/18]

PPE to be used against category II risks

EN388: 2016



3111A

is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN ISO 21420:2020, EN 388:2016 +A1:2018 and is identical to the PPE which is subject to the EU-Type examination (Module B, Annex V of the Regulation), under certificate number 032/2022/0428.02, issued by the Notified Body:

CENTEXBEL (0493)
TECHNOLOGIEPARK 70
B-9052 ZWIJNAARDE
BELGIUM

Guido Van Duren

Director - Regulatory affairs

Ansell

Place: Brussels Date: 2022/03/18

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:

Nitrotough™ N230Y

Products manufactured as of: [2022/03/18]

PPE to be used against category II risks

EN388: 2016



4111A

is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN 388:2016 +A1:2018, EN ISO 21420:2020 and is identical to the PPE which is subject to the EU-Type examination (Module B, Annex V of the Regulation), under certificate number 032/2022/0428, issued by the Notified Body:

CENTEXBEL (0493) TECHNOLOGIEPARK 70 B-9052 ZWIJNAARDE BELGIUM

and is subject to the procedure set out in Annex VI (Module C) of the Regulation.

Guido Van Duren

Director - Regulatory affairs

Ansell

Place: Brussels Date: 2022/03/18

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:

Nitrotough™ N230Y

Products manufactured as of: [2020/01/20] and till: [2022/03/17]

PPE to be used against category II risks



4111A

is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN 388:2016, EN 420:2003 + A1:2009 and is identical to the PPE which is subject to the EU-Type examination (Module B, Annex V of the Regulation), under certificate number 032/2019/1450, issued by the Notified Body:

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

and is subject to the procedure set out in Annex VI (Module C) of the Regulation.

Guido Van Duren

Director - Regulatory affairs

Ansell

Place: Brussels Date: 2020/01/20

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

and authorized representative:
MARIGOLD INDUSTRIAL PORTUGAL
ZONA INDUSTRIAL
APARTADO 41
VILLA NOVA DE POIARES
3350-214 PORTUGAL

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

N230Y

Products manufactured till: [2020/01/19]

PPE to be used against category II risks



is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN 388:2003, EN 420:2003 + A1:2009 and is identical to the PPE which is subject to the EC Type examination; under certificate number 1184 issued by the Notified Body:

INSPEC CERTIFICATION LIMITED (0194)
UPPER WINGBURY COURTTARD, WINGRAVE, AYLESBURY
BUCKINGHANSHIRE, HP22 4LW, UK

and is subject to the procedure set out in Annex VI (Module C) of the Regulation.

Guido Van Duren

Director - Regulatory affairs

Ansell

Place: Brussels Date: 2004/02/26