EU 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® 97-120

Products manufactured as of: [2018/06/25]

PPE to be used against category III risks



4242BP



X1XXXX

is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN 388:2016, EN 407:2004, 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 032/2018/1145, issued by the Notified Body:

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

and is subject to the procedure set out in Annex VII (Module C2) of the Regulation under the supervision of the Notified Body:

Guido Van Duren

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

Director - Regulatory affairs

Ansell

Place: Brussels Date: 2018/06/25

EU 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® Oilfield Services Impact 97-120

Products manufactured till: [2018/06/24]

PPE to be used against category III risks







is in conformity with the provisions of Regulation (EU) 2016/425 and with the standards EN 388:2003, EN 407:2004, EN 420:2003 + A1:2009 and is identical to the PPE which is subject to the EC Type examination; under certificate number 032/2014/1808 issued by the Notified Body:

CENTEXBEL (0493) TECHNOLOGIEPARK 70 B-9052 ZWIJNAARDE BELGIUM

and is subject to the procedure set out in Annex VII (Module C2) of the Regulation under the supervision of the Notified Body:

Guido Van Duren

CENTEXBEL (0493) TECHNOLOGIEPARK 70 B-9052 ZWIJNAARDE

Director - Regulatory affairs

Ansell

Place: Brussels Date: 2014/12/24