EU DECLARATION OF CONFORMITY

FOR PERSONAL PROTECTIVE EQUIPMENT

Originator: J.F ROBLES	**Revision:** 9	**Revision date:** 19.02.2020	**Validity date:** 24.08.2023

PRODUCT	SHIELDskin XTREME™ Sterile Latex 300 DI
DESCRIPTION	DI Powder Free DI washed Pair packed Hand-specific 30cm Cleanroom Gloves
CLASSIFICATION	Personal Protective Equipment (PPE) Category III (Complex Design)

SHIELD Scientific codes	Sizes
69 5551	5.5
69 5552	6
69 5553	6.5
69 5554	7
69 5555	7.5
69 5556	8
69 5557	8.5
69 5558	9
69 5559	10

The manufacturer established in the Union:

SHIELD Scientific B.V.

Dr Willem Dreeslaan 1 – 6721 ND BENNEKOM – THE NETHERLANDS

declares under his/her sole responsibility that the PPE (product codes as mentioned above) described hereafter:

SHIELDskin XTREME™ Sterile Latex 300 DI

is in conformity with the provisions of Regulation (EU) 2016/425 and with the harmonized standards EN ISO 374-1:2016 (as a Type B glove against reagents: K, P & T), EN 374-2:2014 (performance level 3, including protection against viruses), EN 16523-1:2015, EN 374-4:2013, EN ISO 374-5:2016 and EN 420:2003 + A1:2009. This device is identical to the PPE, which is the subject of EU Type Examination (Module B) certificate of conformity no. FI18/961951 issued by the Notified Body:

SGS FIMKO OY (Notified Body No: 0598)
Takomotie 8, FI-00380 Helsinki, Finland

This device is subject to the procedure set out in Article VIII (Module D) of the Regulation under the surveillance of the Notified Body:

SGS FIMKO OY (Notified Body No: 0598)
Takomotie 8, FI-00380 Helsinki, Finland

Signed for and behalf of SHIELD Scientific B.V

J.F ROBLES
General Manager

Date: 19th February 2020
Place: Bennekom
Validity of this declaration: 19th February 2020 until 24th August 2023