

DECLARATION OF CONFORMITY

This UK Declaration of Conformity is issued under the sole responsibility of Ansell Healthcare Europe NV.

Manufacturer Name/Address: Ansell Healthcare Europe NV
Boulevard International 55,
Brussels,
B-1070,
Belgium

UK Responsible Person: Ansell (U.K.) Limited
Building C,
Willerby Hill Business Park,
Willerby, Hull
HU10 6FE
United Kingdom

Risk Class: Class I

GMDN Code & Definition: 56286 - Nitrile examination/treatment glove, non-powdered, non-antimicrobial

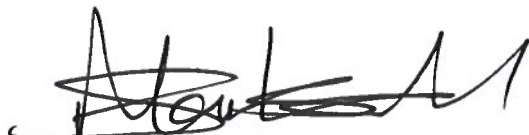
Product Name(s):

Product Name	Size	External Reference Code
MICROFLEX® TQ-601 Soft White Nitrile	XS	TQ-601-XS
	S	TQ-601-S
	M	TQ-601-M
	L	TQ-601-L
	XL	TQ-601-XL

Conformity Assessment Procedure: Part II – Annex VII

We hereby declare that the medical device(s) specified above meet the provision of The Medical Devices Regulations 2002 (UK MDR 2002).

Signed on behalf of the Manufacturer



Name: Samantha Marshall
Position: Director Regulatory Affairs Medical EMEA / APAC
Date: 25 May 2023
Revision: MED\UKMDR\MFXTQ-601SWN\001

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