Ansell Healthcare Europe NV/SA
—
Riverside Business Park – Block J
Boulevard International 55
1070 Brussels, Belgium

T. + 32 (0)2 528 74 00
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Email: info.europe@ansell.com
www.ansell.com

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® 93-833	XS	93833060
	S	93833070
	M	93833080
	L	93833090
	XL	93833100
MICROFLEX® XCEED® XC-INT	XS	XC-INT-XS
	S	XC-INT-S
	M	XC-INT-M
	L	XC-INT-L
	XL	XC-INT-XL

Conformity Assessment Procedure: Part II – Annex VII

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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

Ansell Healthcare Europe NV
Riverside Business Park - Block J
Bld Internationalelaan 55
B-1070 Brussels
BELGIUM

Name: Samantha Marshall
Position: Director Regulatory Affairs Medical EMEA / APAC
Date: 25 May 2023
Revision: MED\UKMDR\MFX93833XC\001