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

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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® LifeStar EC™ 93-868	S	93868070
	M	93868080
	L	93868090
	XL	93868100
	XXL	93868110
	XXXL	93868120
	M-XL	93868000-SAMP

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).

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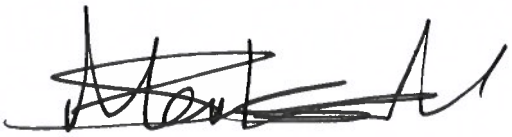
Signed on behalf of the Manufacturer



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BELGIUM

Name: Samantha Marshall
Position: Director Regulatory Affairs Medical EMEA / APAC
Date: 25 May 2023
Revision: MED\UKMDR\MFXLIFST93868\001