

## **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

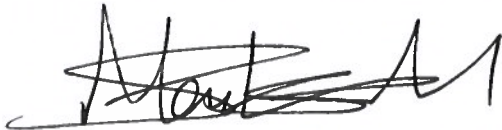
**Product Name(s):**

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

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  
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Position: Director Regulatory Affairs Medical EMEA / APAC  
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