

Ansell Healthcare Europe NV/SA

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Riverside Business Park – Block J
Boulevard International 55
1070 Brussels, Belgium

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Email: info.europe@ansell.com
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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 IIa

#GMDN Code & Definition: 47178 – Hevea-latex surgical glove, non-powdered, non-antimicrobial

Product Name(s):

Product Name	Size	External Reference Code
Medi-Grip® Latex Standard	5.5	330200755
	6.0	330200760
	6.5	330200765
	7.0	330200770
	7.5	330200775
	8.0	330200780
	8.5	330200785
	9.0	330200790

Product Name	Size	External Reference Code
Medi-Grip® PF	5.5	330206055
	6.0	330206060
	6.5	330206065
	7.0	330206070
	7.5	330206075
	8.0	330206080
	8.5	330206085
	9.0	330206090

Product Name	Size	External Reference Code
Medi-Grip® Latex Powder Free	5.5	330200955
	6.0	330200960
	6.5	330200965
	7.0	330200970
	7.5	330200975
	8.0	330200980
	8.5	330200985
	9.0	330200990
	5.5	354101
	6.0	354102
	6.5	354103
	7.0	354104
	7.5	354105
	8.0	354106
	8.5	354107
	9.0	354108

Conformity Assessment Procedure: Part II – Annex V

UKCA Certificate No. 762590

Certified through the British Standards Institution, Approved Body Number 0086.

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

A handwritten signature in black ink, appearing to read "S. Marshall", written over a horizontal line.

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: 24 March 2022
Revision: MED\UKMDR\MEDIGRIPNRLSURG\001

