

EU DECLARATION OF CONFORMITY

This EU 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

SRN Number: BE-MF-000000691

Risk Class: Class IIa

Intended Purpose: A sterile medical device intended as a surgical glove and a protective barrier when worn on the hands of healthcare providers at the surgical site. It is used mainly as a two-way barrier to protect patient and staff from microorganisms and risk of allergy to latex. This is a single-use device.

EMDN Code and Description: T01010201 - Polychloroprene Surgical Gloves

Basic UDI DI: 5414566 GNL340851 NV

Product Name(s):

Product Name	Product Code	Size	Region(s)
Gammex® Non-Latex	340006055	5.5	EMEA/ANZ/APAC
Gammex® Non-Latex	340006060	6	EMEA/ANZ/APAC
Gammex® Non-Latex	340006065	6.5	EMEA/ANZ/APAC
Gammex® Non-Latex	340006070	7	EMEA/ANZ/APAC
Gammex® Non-Latex	340006075	7.5	EMEA/ANZ/APAC
Gammex® Non-Latex	340006080	8	EMEA/ANZ/APAC
Gammex® Non-Latex	340006085	8.5	EMEA/ANZ/APAC
Gammex® Non-Latex	340006090	9	EMEA/ANZ/APAC
Gammex® Non-Latex	8511	5.5	NA/India
Gammex® Non-Latex	8512	6	NA/India
Gammex® Non-Latex	8513	6.5	NA/India
Gammex® Non-Latex	8514	7	NA/India

Product Name	Product Code	Size	Region(s)
Gammex® Non-Latex	8515	7.5	NA/India
Gammex® Non-Latex	8516	8	NA/India
Gammex® Non-Latex	8517	8.5	NA/India
Gammex® Non-Latex	8518	9	NA/India

Conformity Assessment Procedure: Annex IX.

CE Certificate No: MDR 763361.

Certified through the British Standards Institution, Notified Body Number 2797.

We hereby declare that the medical device(s) specified above meet the provision of the Regulation (EU) MDR 2017/745 for medical devices.

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
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