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Product Certificate Thermo Scientific Nalgene and Nunc Products

Thermo Fisher Scientific hereby certifies that the product identified below is produced, inspected and found to be in compliance with product and quality specification requirements as documented in our ISO 13485:2003 Quality Management System (QMI-SAI Global File No. 1606319 and 1606321) in the USA.

Product Compliance Manager

Certificate issued: 06/07/2016

The following information represents Product Certification for: Item#: 374223-BR

Description: Nunc Cryobank 2.0mL, Non-2D, w/Cap, BC Rk Lot#: 1177867

Use Before: 05/28/2021

Manufactured:

05/23/2016

Part Number	Description	Common Name	DMF#	Cytotoxicity	USP Class VI	FDA Compliance - 21 CFR
1-0025-70	VIAL,CRYOBANK,2.0mL,W/BC HEAD	COMPONENT PART				
8-0071-22	Resin, PP, Black, INJ, Cryobank	COMPONENT PART				
14117MR	RESIN,PP,HIMONT PF-511	PP, HOMO, NATURAL, INJ.	26106	PASSED	PASSED	177.1520(a)(1)(i) and (c)1.1a, 177.1520(b), use conditions A-H
14288MR	COLORANT,BLACK,PP	COLORANT, BLACK, PP, INJ.	N/A	N/A	N/A	N/A
8-0028-04	RESIN,PPCO,RAD STER,INJ	POLYPROPYLENE COPOLYMER	7478	PASSED	PASSED	177.1520 (a)(3)(i) & (c)3.1(a)except for cooking, (useconditions C-H)
1-0025-73	CAP,CRYOVIAL 2/5mL NON-PIERCEABLE	COMPONENT PART				
8-0028-04	RESIN,PPCO,RAD STER,INJ	POLYPROPYLENE COPOLYMER	7478	PASSED	PASSED	177.1520 (a)(3)(i) & (c)3.1(a)except for cooking, (useconditions C-H)
8-0005-22	RESIN,TPV,NAT,45 DURO SANTOPRENE	THERMOPLASTIC RUBBER	12719	PASSED	PASSED	N/A

If N/A appears in any of the columns above it means the information is not available. Any item listed as "COMPONENT PART" will show blank in the DMF#, Cytotoxicity, USP Class VI, and FDA Compliance Information columns.

If the word "PASSED" appears in the USP Class VI column next to the resin listing, this material has passed USP Class VI requirements, latest Volume, as part of our initial test approval protocol.

If the word "PASSED" appears in the Cytotoxicity column next to the resin listing, this material was tested and shown to be non-cytotoxic as part of our initial test approval protocol, using either mouse fibroblast L929 cells or the more sensitive human diploid lung cell lines WI-38 or MRC-5.