ThermoFisher SCIENTIFIC

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Product Certificate

Thermo Scientific
Nalgene and Nunc Products

Thermo Fisher Scientific hereby certifies that the product identified below is manufactured and/or distributed according to the requirements of product and quality specifications as maintained in our quality management system which is compliant to ISO 13485:2016 (BSI Certificate Number: FM 653694) in the USA.

Alan E. Hatch
Sr. Quality Manager

The following information represents Product Certification for: Item#: **3110-42**

Description: **BIOTAIN,PETG,1L,ST,LP** Lot#: **1374644** Use Before: **12/19/2027** Manufactured: **12/20/2022**

Part Number	Description	Common Name	DMF#	Cytotoxicity	USP Class VI	FDA Compliance - 21 CFR
1-0449-91	CLOS,48MM,Q/A,W/O SLRNG,PP,WHT	COMPONENT PART				
8-0028-16P	RESIN,PPCO,RAD STAB,WHITE,INJ	COLOR MIX (PPCO, RAD STAB,WHT)	N/A	PASSED	PASSED	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)
8-0099-34	COLOR,WHT,MULTI	COLORANT, WHITE	16513	PASSED	PASSED	177.1350, 1520, 1620,178.3297, 181.28
1-0449-77P	BIOTAINER,1L,SQ,Q/A,PETG	COMPONENT PART				
8-0001-32	RESIN,PETG,IBM,EBM,INJ	COPOLYESTER, PETG NATURAL	9987	PASSED	PASSED	177.1315(b)(1) and 174.5

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.

Product was Gamma Irradiation Sterilized. Product was dosimetric released per ANSI/AAMI/ISO 11137 guidelines. Product was determined to be non-pyrogenic at a level < 0.5 EU/ml per USP < 85 > .

The foam liner and PP closure materials have been tested and shown to comply with USP Class VI requirements and also have been shown to be non-cytotoxic. The liner material meets requirements of CFR 21, Sections 175.300, 177.1520, 177.1210, 178.2010 of the Federal Food and Drug Act. The closure materials meet requirements of CFR 21, Section 177.1520 of the Federal Food and Drug Act.

ANIMAL DERIVED MATERIALS (BSE/TSE) - All resins and colorants used at the manufacturing site comply with the latest revision of EMA/410/01 section 6.4. The purchased component(s) used in this product also comply with the latest revision of EMA/410/01 section 6.4.