

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.



Janet Cosgrove
Raw Materials and
Product Compliance Manager

The following information represents Product Certification for: Item#: **5000-0020**

Certificate issued: **11/07/2016**

Description: **CRYOGENIC VIAL 2.0ML, PPCO**

Lot#: **1188080**

Use Before: **11/28/2021**

Manufactured: **11/01/2016**

| Part Number | Description | Common Name | DMF# | Cytotoxicity | USP Class VI | FDA Compliance - 21 CFR |
|-------------|-------------------------------|---------------------------|------|--------------|--------------|---|
| 1-3825-83 | CLOS,CRYOVL,9X9,HDPE,BULK PK | COMPONENT PART | | | | |
| 8-0042-02 | RESIN,HDPE,HF,INJ | HIGH DENSITY POLYETHYLENE | 1646 | PASSED | PASSED | 176.170(c), 177.1520(c)3.2a |
| 1-3825-92 | VIAL,CRYO,2.0ML,2-1/2 TURN,PP | 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) |

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 > .

Product is certified to be free of detectable RNase/DNase contamination. This test is performed using the nuclease assay method with a detection limit of 8×10^{-7} Kunitz unit/ul for DNase and 1.9×10^{-10} Kunitz unit/ul for RNase.