



## CERTIFICATE OF CONFORMANCE

**Product Part Number:** PEN-12IR  
**Lot Number:** 054629-000  
**Irradiation Reference Number:** 0012114-00  
**Date of manufacture:** 04/27/2026  
**Manufacturing/Processing Site:** MICRONOVA MANUFACTURING LLC  
3431 WEST LOMITA BLVD. TORRANCE, CA 90505  
**Irradiation Site:** STERIS ISOMEDIX  
1000 SARAH PL. ONTARIO, CA 91761  
**Shelf Life:** 24 MONTHS\*  
**Description:** CLEANROOM PEN, CLICK, FINE POINT, BLUE, IRR'D  
(100/CS)  
**Size -**  
**Width:** N/A  
**Length:** N/A  
**Height:** N/A

*This certification is provided as full assurance that the following product code and lot number was manufactured in accordance with prescribed procedures and specifications.*

5/4/2026

**Authorized Signature**  
**Quality Control Department**

**Date**

**CC:** With Product  
Customer File  
C of C File

**\*Refer to Certificate of Processing for irradiation complete date**

3431 WEST LOMITA BOULEVARD • TORRANCE, CA 90505-5010 • 301 LEORA LANE (BUILDING 4, SUITE 420), THE COLONY, TX 75056  
Tel. (310) 784-6990 • FAX (310) 784-6980

# STERIS: Certificate Of Processing

Prepared For MICRONOVA MFG INC (2727)

Gamma Process Run ID 1113-30617A

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<u>Product Code</u>	<u>Lot Number</u>	<u>Quantity</u>	<u>UOM</u>
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PEN-12IR	054629-000	80	Case
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Processing Run Start Date 30-Apr-2026 11:04 PM

Processing Run End Date 01-May-2026 4:44 AM

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Minimum Specified Dose (kGy)	25.0	Minimum Delivered Dose (kGy)	28.5
Maximum Specified Dose (kGy)	45.0	Maximum Delivered Dose (kGy)	38.5

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PO# 0012114-00

Product meets Customer specifications; zero nonconformities occurred during this irradiation run.

**Gamma Process Run Approval authorized by STERIS**

DateTime Esigned 01-May-2026 8:45 AM

Operating facilities are in compliance with applicable local/state and federal regulations providing services under a certified quality system which meets the following requirements (if applicable): FDA QMSR, ISO 9001, ISO 13485 current certified version, national pharmaceutical GMP (EU GMP, Commission Directive 2017/1572, Veterinary Regulation (EU) 2019/6, etc.) and is in alignment with ISO 11137 current certified version. STERIS AST certifies that these processed items received the indicated doses within the precision and accuracy of the dosimetry system used. The product has been processed as per the requirements of the Customer Agreement and/or Customer Specification.

## Processing Location

1000 Sarah Place  
Ontario CA 91761  
United States

**APPROVED**

*By Jenicah Baquir at 8:56 am, May 01, 2026*

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