

# **CERTIFICATE OF CONFORMANCE**

| Product Part Number: CCS           | C-6IR                                   |
|------------------------------------|---|
| Lot Number: 0503                   | 108-000                                 |
| Irradiation Reference Number: 0010 | 0711-00                                 |
| Lot Quantity: 30 C                 | ASE(S)                                  |
| Date of manufacture: 06/1          | .0/2025*                                |
| Shelf Life: 60 N                   | IONTHS                                  |
| Description: CUR                   | TAIN CLEANER, NOVAPOLY, IRR'D (12PR/CS) |
| Size -                             |   |
| Width: 2 ½"                        |   |
| Length: 14 ½                       | , <sup>1</sup>                          |
| 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.

### 6/20/2025

## Authorized Signature Quality Control Department

Date

CC: With Product Customer File C of C File

\*Refer to Certificate of Processing for irradiation complete date

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## **STERIS: Gamma Certificate Of Processing**

**MICRONOVA MFG INC (2727) Prepared For** Gamma Process Run ID 1113-26354A Product Code Lot Number Quantity UOM CCSC-6IR 050108-000 30 Case Processing Run Start Date 20-Jun-2025 2:08 AM Processing Run End Date 20-Jun-2025 9:12 AM 29.1 Minimum Specified Dose (kGy) 25.0 Minimum Delivered Dose (kGy) Maximum Specified Dose (kGy) 40.0 Maximum Delivered Dose (kGy) 38.7

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

#### **Other Information**

0010711-00

## Gamma Process Run Approval authorized by STERIS

DateTime Esigned 20-Jun-2025 10:25 AM

Operating facilities are in compliance with applicable state and federal regulations (FDA, NRC, EPA, and OSHA) and provide services under a quality system which meets the requirements of FDA QSR, EN/ISO 13485:2016, and in alignment with EN ANSI/AAMI/ISO 11137:2017. STERIS certifies that these processed items received the indicated doses within the precision and accuracy of the dosimetry system used.

Processing Location

1000 Sarah Place Ontario CA 91761 United States

> **APPROVED** By Jenicah Baquir at 10:27 am, Jun 20, 2025

> > Document ID: 226184