

K240429 Trilogy Tissue MarkerNov 7, 2024
268 days to decisionK240429 · Product code: **NEU** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k240429/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Marker, Radiographic, Implantable (NEU)
Date received	Feb 13, 2024
Decision date	Nov 7, 2024
Days to decision	268 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Inrad, Inc.
Location	Grand Rapids, MI, US
Contact	Heidi Halverson
510(k) history	4 submissions · 4 cleared · 2011-2025

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k240429/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 14, 2026