

**K942489 ELECTROSURGICAL CUTTING & COAGULATION
DEVICE**Jul 25, 1994
61 days to decisionK942489 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k942489/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	May 25, 1994
Decision date	Jul 25, 1994
Days to decision	61 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Megadyne Medical Products, Inc.
Location	Murray, UT, US
Contact	DREW D WEAVER
510(k) history	50 submissions · 48 cleared · 1990-2024

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k942489/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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