

K072559 SUCTION COAGULATOROct 5, 2007
24 days to decisionK072559 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k072559/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Sep 11, 2007
Decision date	Oct 5, 2007
Days to decision	24 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k072559/> 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