

K992045 ELECTROSURGICAL ELECTRODE FAMILYJan 27, 2000
224 days to decisionK992045 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k992045/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jun 17, 1999
Decision date	Jan 27, 2000
Days to decision	224 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Genesis Medical, Inc.
Location	Sunnyvale, CA, US
Contact	THOMAS WEHMAN
510(k) history	1 submissions · 1 cleared · 2000-2000

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k992045/> 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 July 3, 2026