

**K001151 ELECTROSURGICAL ELECTRODE FAMILY**Jun 27, 2000  
78 days to decisionK001151 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k001151/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Apr 10, 2000
Decision date	Jun 27, 2000
Days to decision	78 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Novasys Medical, Inc.</b>
Location	Sunnyvale, CA, US
Contact	THOMAS WEHMAN
510(k) history	5 submissions · 5 cleared · 2000-2004

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k001151/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 30, 2026