

**K013139 STELLARTECH COAGULATION SYSTEM**Dec 18, 2001  
90 days to decisionK013139 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k013139/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Sep 19, 2001
Decision date	Dec 18, 2001
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Stellartech Research Corp.</b>
Location	Sunnyvale, CA, US
Contact	JAMES R SANTOS
510(k) history	11 submissions · 11 cleared · 2000-2006

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k013139/> 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 29, 2026