

**K023311 LOOP MICROWAVE ABLATION PROBE**Apr 9, 2003  
188 days to decisionK023311 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k023311/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Oct 3, 2002
Decision date	Apr 9, 2003
Days to decision	188 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Vivant Medical, Inc.</b>
Location	Portola Valley, CA, US
Contact	STEVEN KIM
510(k) history	10 submissions · 10 cleared · 2000-2005

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k023311/> 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