

**K050223 VIVAWAVE MICROWAVE ABLATION SYSTEM,
COAXIAL INTRODUCER**Feb 24, 2005
24 days to decisionK050223 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k050223/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jan 31, 2005
Decision date	Feb 24, 2005
Days to decision	24 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Vivant Medical, Inc.
Location	Portola Valley, CA, US
Contact	KRISTINE FOSS
510(k) history	10 submissions · 10 cleared · 2000-2005

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k050223/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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