

**K180584 Viveve RF System, Secure**Apr 6, 2018  
32 days to decisionK180584 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k180584/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Mar 5, 2018
Decision date	Apr 6, 2018
Days to decision	32 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Viveve, Inc.</b>
Location	Sunnyvale, CA, US
Contact	Suzon Lommel
510(k) history	3 submissions · 3 cleared · 2016-2022

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k180584/> 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 May 27, 2026