

**K072493 ENSEAL TISSUE SEALING DEVICE**Sep 27, 2007  
22 days to decisionK072493 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k072493/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Sep 5, 2007
Decision date	Sep 27, 2007
Days to decision	22 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Surgrx, Inc.</b>
Location	Mountain View, CA, US
Contact	LINDA OLESON
510(k) history	12 submissions · 12 cleared · 2003-2008

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k072493/> 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 24, 2026