

**K081431 MEG LAPAROSCOPIC ELECTRODES**Jul 24, 2008  
64 days to decisionK081431 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k081431/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	May 21, 2008
Decision date	Jul 24, 2008
Days to decision	64 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Bovie Medical</b>
Location	St. Petersburg, FL, US
Contact	RICHARD KOZLOFF
510(k) history	8 submissions · 8 cleared · 2006-2010

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k081431/> 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 21, 2026