

**K981188 ELECTROSURGICAL CAUTERY PROBES**May 20, 1998  
48 days to decisionK981188 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k981188/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Apr 2, 1998
Decision date	May 20, 1998
Days to decision	48 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Microline Pentax, Inc.</b>
Location	Devers, MA, US
Contact	JACQUELINE E MASSE
510(k) history	12 submissions · 12 cleared · 1988-2002

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k981188/> 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