

**K962119 RE-NEW LAPAROSCOPIC INSTRUMENTS**Aug 9, 1996  
70 days to decisionK962119 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k962119/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	May 31, 1996
Decision date	Aug 9, 1996
Days to decision	70 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Microline Pentax, Inc.</b>
Location	Devers, MA, US
Contact	HUGHES DE LAFORCADE
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/k962119/> 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