

**K081559 GENESIS**Aug 28, 2008  
86 days to decisionK081559 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k081559/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jun 3, 2008
Decision date	Aug 28, 2008
Days to decision	86 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Medisiss</b>
Location	Sisters, OR, US
Contact	BRANDI JAMES
510(k) history	6 submissions · 6 cleared · 2003-2011

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k081559/> 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 18, 2026