

**K120431 HALO90 ULTRA ABLATION CATHETER**May 3, 2012  
80 days to decisionK120431 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k120431/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Feb 13, 2012
Decision date	May 3, 2012
Days to decision	80 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Barrx Medical, Inc.</b>
Location	Sunnyvale, CA, US
Contact	DAWN CHANG
510(k) history	3 submissions · 3 cleared · 2010-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k120431/> 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 14, 2026