

**K071543 HALO360's; COAGULATION CATHETER**Jun 29, 2007  
24 days to decisionK071543 · Product code: **GEI** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k071543/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrosurgical, Cutting & Coagulation & Accessories (GEI)
Date received	Jun 5, 2007
Decision date	Jun 29, 2007
Days to decision	24 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Barrx Medical, Incorporated</b>
Location	Washington, Dc, DC, US
Contact	VIORICA FILIMON
510(k) history	13 submissions · 13 cleared · 2005-2010

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k071543/> 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