

**K031001 ACMI VISTA CTR BIPOLAR LOOP ELECTRODE**May 21, 2003  
51 days to decisionK031001 · Product code: **FAS** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k031001/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrode, Electrosurgical, Active, Urological (FAS)
Date received	Mar 31, 2003
Decision date	May 21, 2003
Days to decision	51 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Acmi Corporation</b>
Location	Southborough, MA, US
Contact	GABRIEL J MURACA
510(k) history	16 submissions · 16 cleared · 2002-2006

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k031001/> 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 29, 2026