

**K060573 BURBANK TENACULUM**Apr 21, 2006  
46 days to decisionK060573 · Product code: **HDC** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k060573/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tenaculum, Uterine (HDC)
Date received	Mar 6, 2006
Decision date	Apr 21, 2006
Days to decision	46 days
Third-party review	Yes
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Vascular Control Systems, Inc.</b>
Location	San Juan Capistrano, CA, US
Contact	KATHLEEN ROBERTS
510(k) history	9 submissions · 9 cleared · 2002-2006

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k060573/> 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