

**K013683 MODIFICATION TO: ABBOTT VASCULAR SUTURE ANASTOMOSIS DEVICE**Dec 6, 2001  
29 days to decisionK013683 · Product code: **GAW** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k013683/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Suture, Nonabsorbable, Synthetic, Polypropylene (GAW)
Date received	Nov 7, 2001
Decision date	Dec 6, 2001
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Abbott Vascular, Inc.</b>
Location	Redwood, CA, US
Contact	PATTY HEVEY
510(k) history	20 submissions · 17 cleared · 2000-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k013683/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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