

**K864401 PLURIMUS CARDIAC EVALUATION SYSTEM**Mar 3, 1987  
117 days to decisionK864401 · Product code: **DSI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k864401/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Nov 6, 1986
Decision date	Mar 3, 1987
Days to decision	117 days
Third-party review	No

**APPLICANT**

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Company	<b>Vivix Corp.</b>
Location	San Jose, CA, US
Contact	L. C BADAGLIACCA
510(k) history	1 submissions · 1 cleared · 1987-1987

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k864401/>, 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 July 7, 2026