

**K873041 A-V RECORDER**Feb 4, 1988  
184 days to decisionK873041 · Product code: **DPT** · CardiovascularSource: <https://www.510kdatabase.net/k873041/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Probe, Blood-flow, Extravascular (DPT)
Date received	Aug 4, 1987
Decision date	Feb 4, 1988
Days to decision	184 days
Third-party review	No

**APPLICANT**

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Company	<b>Life Sciences Manufacturing, Inc.</b>
Location	Mchenry, IL, US
Contact	JAMES H VEZINA
510(k) history	12 submissions · 12 cleared · 1979-1991

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k873041/>, 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