

**K842133 EYRTHROCATH DOUBLE-LUMEN CENTRAL VENOUS**Feb 19, 1985  
265 days to decisionK842133 · Product code: **FOZ** · General HospitalSource: <https://www.510kdatabase.net/k842133/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Short-term Less Than 30 Days (FOZ)
Date received	May 30, 1984
Decision date	Feb 19, 1985
Days to decision	265 days
Third-party review	No

**APPLICANT**

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Company	<b>Cardio Search, Inc.</b>
Location	Walker, MI, US
Contact	LARRY H BRIM
510(k) history	4 submissions · 4 cleared · 1983-1985

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k842133/>; 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 8, 2026