

**K831645 TEMP. BIPOLAR EPICARDIAL LEAD**Jul 26, 1983  
64 days to decisionK831645 · Product code: **LDF** · CardiovascularSource: <https://www.510kdatabase.net/k831645/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	May 23, 1983
Decision date	Jul 26, 1983
Days to decision	64 days
Third-party review	No

**APPLICANT**

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Company	<b>Cordic Corp.</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1983-1983

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k831645/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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