

**K882045 HIGH TORQUE TEMPORARY PERVENOUS LEAD  
W/DEPTH MARK**Jul 12, 1988  
57 days to decisionK882045 · Product code: **LDF** · Cardiovascular  
Source: <https://www.510kdatabase.net/k882045/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	May 16, 1988
Decision date	Jul 12, 1988
Days to decision	57 days
Third-party review	No

**APPLICANT**

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Company	<b>Cordis Leads, Inc.</b>
Location	Suffield, CT, US
Contact	DUANE A SCHULTZ
510(k) history	4 submissions · 4 cleared · 1988-1988

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k882045/>; 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