

**K944957 TEMPORARY MYOCARDIAL PACING LEAD, MODEL 6500**Jul 14, 1995  
280 days to decisionK944957 · Product code: **LDF** · Cardiovascular  
Source: <https://www.510kdatabase.net/k944957/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	Oct 7, 1994
Decision date	Jul 14, 1995
Days to decision	280 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Medtronic Blood Systems, Inc.</b>
Location	Anaheim, CA, US
Contact	DIANA SALDITT
510(k) history	15 submissions · 15 cleared · 1988-1995

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k944957/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://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](https://accessdata.fda.gov). - Generated May 14, 2026