

**K922269 UNIPOLAR TEMPORARY MYOCARDIAL PACING
WIRE, 6493**Nov 23, 1992
193 days to decisionK922269 · Product code: **LDF** · Cardiovascular
Source: <https://www.510kdatabase.net/k922269/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	May 14, 1992
Decision date	Nov 23, 1992
Days to decision	193 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	The Heart Valve Div. Medtronic Cardiovascular Surg
Location	Minneapolis, MN, US
Contact	DIANA K SALDITT
510(k) history	5 submissions · 4 cleared · 1991-1998

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k922269/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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