

K012452 MODEL 6500 UNIPOLAR TEMPORARY MYOCARDIAL PACING LEADAug 14, 2001
13 days to decisionK012452 · Product code: **LDF** · Cardiovascular
Source: <https://www.510kdatabase.net/k012452/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	Aug 1, 2001
Decision date	Aug 14, 2001
Days to decision	13 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Medtronic Vascular
Location	Walker, MI, US
Contact	TINA BENOIT
510(k) history	475 submissions · 453 cleared · 1977-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k012452/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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