

**K954809 UNIPOLAR TEMPORARY MYOCARDIAL PACING WIRE
MODEL 6494**Dec 13, 1995
55 days to decisionK954809 · Product code: **LDF** · Cardiovascular
Source: <https://www.510kdatabase.net/k954809/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	Oct 19, 1995
Decision date	Dec 13, 1995
Days to decision	55 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k954809/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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