

**K850622 OSCOR MEDICAL MODEL TME 64S BIPOLAR
TEMP/HEART WIR**Apr 10, 1985
50 days to decisionK850622 · Product code: **LDF** · Cardiovascular
Source: <https://www.510kdatabase.net/k850622/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	Feb 19, 1985
Decision date	Apr 10, 1985
Days to decision	50 days
Third-party review	No

APPLICANT

Company	Oscor Medical Corp.
Location	Washington, DC, US
Contact	ALLEN M FOX
510(k) history	31 submissions · 30 cleared · 1985-1997

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k850622/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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