

**K884866 TEMP. PACING LEADS W/DEPTH MARKINGS TA-1
AND TB-1**Mar 22, 1989
121 days to decisionK884866 · Product code: **LDF** · Cardiovascular
Source: <https://www.510kdatabase.net/k884866/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	Nov 21, 1988
Decision date	Mar 22, 1989
Days to decision	121 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k884866/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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