

K830772 TEMP. PACING LEAD, BIPOLAR #TH60, 90/110Apr 18, 1983
38 days to decisionK830772 · Product code: **LDF** · CardiovascularSource: <https://www.510kdatabase.net/k830772/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	Mar 11, 1983
Decision date	Apr 18, 1983
Days to decision	38 days
Third-party review	No

APPLICANT

Company	Oscor, Inc.
Location	Palm Harbor, FL, US
510(k) history	49 submissions · 46 cleared · 1979-2021

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k830772/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 27, 2026