

K831695 TEMP. CARDIAC PACING WIRE TME 60/61/65Oct 20, 1983
148 days to decisionK831695 · Product code: **LDF** · Cardiovascular
Source: <https://www.510kdatabase.net/k831695/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	May 25, 1983
Decision date	Oct 20, 1983
Days to decision	148 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k831695/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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