FDA 510(k) Premarket Notification

K831804 MULTIPOLAR TEMPORARY CARDIAC

CLEARED

Oct 20, 1983
139 days to decision

K831804 · Product code: **LDF** · Cardiovascular
Source: https://www.510kdatabase.net/k831804/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	Jun 3, 1983
Decision date	Oct 20, 1983
Days to decision	139 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.net
Device record: https://www.510kdatabase.net/k831804/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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