FDA 510(k) Premarket Notification

K850622 OSCOR MEDICAL MODEL TME 64S BIPOLAR TEMP/HEART WIR

CLEARED

Apr 10, 1985
50 days to decision

K850622 · 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