FDA 510(k) Premarket Notification

K872972 TEMPORARY PACEMAKER ELECTRODE

CLEARED

Dec 3, 1987
127 days to decision

K872972 · Product code: **LDF** · Cardiovascular
Source: https://www.510kdatabase.net/k872972/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	Jul 29, 1987
Decision date	Dec 3, 1987
Days to decision	127 days
Third-party review	No

APPLICANT

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