FDA 510(k) Premarket Notification

K884866 TEMP. PACING LEADS W/DEPTH MARKINGS TA-1 AND TB-1

CLEARED

Mar 22, 1989
121 days to decision

K884866 · Product code: **LDF** · Cardiovascular
Source: https://www.510kdatabase.net/k884866/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	Nov 21, 1988
Decision date	Mar 22, 1989
Days to decision	121 days
Third-party review	No

APPLICANT

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