FDA 510(k) Premarket Notification

K862330 MODIFIED OSCOR PY PACING LEADS

CLEARED

Aug 8, 1986
50 days to decision

K862330 · Product code: **DTB** · Cardiovascular
Source: https://www.510kdatabase.net/k862330/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Permanent Pacemaker Electrode (DTB)
Date received	Jun 19, 1986
Decision date	Aug 8, 1986
Days to decision	50 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