FDA 510(k) Premarket Notification

K882345 PERMANENT PACING LEADS PYBJ SERIES

CLEARED

Oct 28, 1988
143 days to decision

K882345 · Product code: **DTB** · Cardiovascular
Source: https://www.510kdatabase.net/k882345/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Permanent Pacemaker Electrode (DTB)
Date received	Jun 7, 1988
Decision date	Oct 28, 1988
Days to decision	143 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