

K882345 PERMANENT PACING LEADS PYBJ SERIESOct 28, 1988
143 days to decisionK882345 · Product code: **DTB** · CardiovascularSource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k882345/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 3, 2026