

**K041809 PERMANENT PACING LEAD, MODEL PY2**Aug 6, 2004  
31 days to decisionK041809 · Product code: **DTB** · Cardiovascular  
Source: <https://www.510kdatabase.net/k041809/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Permanent Pacemaker Electrode (DTB)
Date received	Jul 6, 2004
Decision date	Aug 6, 2004
Days to decision	31 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>Oscor, Inc.</b>
Location	Palm Harbor, FL, US
Contact	MILA DOSKOCIL
510(k) history	49 submissions · 46 cleared · 1979-2021

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k041809/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 27, 2026