

**K020669 PERMANENT PACING LEAD, MODELS PETITE,
PETITE R, J, RJ, PETITE B, RB, JB, RJB**Mar 28, 2002
27 days to decisionK020669 · Product code: **DTB** · Cardiovascular
Source: <https://www.510kdatabase.net/k020669/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Permanent Pacemaker Electrode (DTB)
Date received	Mar 1, 2002
Decision date	Mar 28, 2002
Days to decision	27 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k020669/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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