

**K830950 VASCULAR PRODUCTS IMPLANTABLE PACING**Apr 18, 1983  
24 days to decisionK830950 · Product code: **DTB** · CardiovascularSource: <https://www.510kdatabase.net/k830950/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Permanent Pacemaker Electrode (DTB)
Date received	Mar 25, 1983
Decision date	Apr 18, 1983
Days to decision	24 days
Third-party review	No

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k830950/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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