

**K811879 MODEL A UNIPOLAR ENDOCARDIAL LEAD**Aug 3, 1981  
33 days to decisionK811879 · Product code: **DTB** · CardiovascularSource: <https://www.510kdatabase.net/k811879/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Permanent Pacemaker Electrode (DTB)
Date received	Jul 1, 1981
Decision date	Aug 3, 1981
Days to decision	33 days
Third-party review	No

**APPLICANT**

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Company	<b>Cook Pacemaker Corp.</b>
Location	Mchenry, IL, US
510(k) history	34 submissions · 31 cleared · 1981-1997

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

Device record: <https://www.510kdatabase.net/k811879/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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