

**K920822 CARDIOVASCULAR PERMANENT PACEMAKER  
ELECTRODE**Jan 22, 1993  
333 days to decisionK920822 · Product code: **DTB** · Cardiovascular  
Source: <https://www.510kdatabase.net/k920822/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - SP
Submission type	Traditional
Device classification	Permanent Pacemaker Electrode (DTB)
Date received	Feb 24, 1992
Decision date	Jan 22, 1993
Days to decision	333 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Siemens-Pacesetter, Inc.</b>
Location	Sylmar, CA, US
Contact	GLENN THOMPSON
510(k) history	7 submissions · 2 cleared · 1992-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k920822/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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