

**K872551 SEAMED MODEL 3200**Jul 31, 1987  
32 days to decisionK872551 · Product code: **DTE** · CardiovascularSource: <https://www.510kdatabase.net/k872551/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pulse-generator, Pacemaker, External (DTE)
Date received	Jun 29, 1987
Decision date	Jul 31, 1987
Days to decision	32 days
Third-party review	No

**APPLICANT**

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Company	<b>Seamed Corp.</b>
Location	Walker, MI, US
Contact	MARCIA A WARRENDER
510(k) history	18 submissions · 18 cleared · 1982-1992

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k872551/>, 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 July 6, 2026