

**K840523 SERVOMED PULSE AMPLIFIER MODULE-206-**Oct 29, 1984  
265 days to decisionK840523 · Product code: **DRT** · CardiovascularSource: <https://www.510kdatabase.net/k840523/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Feb 7, 1984
Decision date	Oct 29, 1984
Days to decision	265 days
Third-party review	No

**APPLICANT**

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Company	<b>Litton Medical Electronics</b>
Location	Walker, MI, US
Contact	CASARSA
510(k) history	38 submissions · 38 cleared · 1982-1985

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k840523/>; 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 May 27, 2026