

**K832250 SERVOMED BASIC BEDSIDE UNIT SMA 102-**Jun 1, 1984  
326 days to decisionK832250 · Product code: **DRT** · CardiovascularSource: <https://www.510kdatabase.net/k832250/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Jul 11, 1983
Decision date	Jun 1, 1984
Days to decision	326 days
Third-party review	No

**APPLICANT**

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Company	<b>Litton Medical Electronics</b>
Location	Walker, MI, US
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/k832250/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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