

**K833000 I-CHANNEL RECORDER**Jun 1, 1984  
288 days to decisionK833000 · Product code: **DSF** · CardiovascularSource: <https://www.510kdatabase.net/k833000/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Recorder, Paper Chart (DSF)
Date received	Aug 18, 1983
Decision date	Jun 1, 1984
Days to decision	288 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/k833000/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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