

**K842715 DOCUMENTOR**Nov 19, 1984  
130 days to decisionK842715 · Product code: **DSI** · CardiovascularSource: <https://www.510kdatabase.net/k842715/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Jul 12, 1984
Decision date	Nov 19, 1984
Days to decision	130 days
Third-party review	No

**APPLICANT**

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Company	<b>Life Technology</b>
Location	Corona, CA, US
Contact	PAT BURRELL
510(k) history	1 submissions · 1 cleared · 1984-1984

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k842715/>; 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 8, 2026