

**K896967 HTS 820 CARDIAC MONITOR**Mar 19, 1990  
91 days to decisionK896967 · Product code: **DSI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k896967/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Dec 18, 1989
Decision date	Mar 19, 1990
Days to decision	91 days
Third-party review	No

**APPLICANT**

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Company	<b>Life Sensing Instrument Company, Inc.</b>
Location	Tullahoma, TN, US
Contact	B. Q CUTSHAW
510(k) history	5 submissions · 5 cleared · 1986-1997

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k896967/>; 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 June 30, 2026