

**K930095 SPACELABS MEDICAL PATIENT MONITOR**Sep 3, 1993  
238 days to decisionK930095 · Product code: **DSI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k930095/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Jan 8, 1993
Decision date	Sep 3, 1993
Days to decision	238 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Spacelabs Medical, Inc.</b>
Location	Hamden, CT, US
Contact	RAYMOND W GIFFORD
510(k) history	35 submissions · 34 cleared · 1993-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k930095/> 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 25, 2026