

**K103142 SPACELABS MULTI-PARAMETER MODULE**Mar 3, 2011  
129 days to decisionK103142 · Product code: **DSI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k103142/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Oct 25, 2010
Decision date	Mar 3, 2011
Days to decision	129 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Spacelabs Healthcare, Inc.</b>
Location	Nederland, CO, US
Contact	THOMAS KROENKE
510(k) history	1 submissions · 1 cleared · 2011-2011

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k103142/> 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 19, 2026