

**K983996 SPACELABS MEDICAL ULTRAVIEW DIGITAL  
TELEMTRY SYSTEM**Jun 3, 1999  
205 days to decisionK983996 · Product code: **MHX** · Cardiovascular  
Source: <https://www.510kdatabase.net/k983996/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient(with Arrhythmia Detection Or Alarms) (MHX)
Date received	Nov 10, 1998
Decision date	Jun 3, 1999
Days to decision	205 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Spacelabs Medical, Inc.</b>
Location	Hamden, CT, US
Contact	NANCY J GERTLAR
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/k983996/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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