

K950779 HEART RATE VARIABILITY SOFTWARE OPTIONApr 25, 1995
63 days to decisionK950779 · Product code: **DPS** · CardiovascularSource: <https://www.510kdatabase.net/k950779/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrocardiograph (DPS)
Date received	Feb 21, 1995
Decision date	Apr 25, 1995
Days to decision	63 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Spacelabs, Inc.
Location	Mchenry, IL, US
Contact	RUSS GARRISON
Website	https://www.spacelabshealthcare.com
510(k) history	46 submissions · 46 cleared · 1976-1996

Spacelabs, Inc. is a medical device company based in McHenry, US. The company specializes in patient monitoring and diagnostic cardiology solutions. Spacelabs received FDA 510(k) clearances from total submissions. The company's regulatory activity spans from 1976 to 1996, with Cardiovascular devices representing the dominant focus of its portfolio. This historical record reflects the company's early contributions to cardiac monitoring and anesthesiology device development. Notable cleared devices include multigas analyzers, capnograph modules, patient care management syst...
