

**K925510 MODEL 90478 & 90479 TELEMETRY REC.
MODULE&HOUSING**Oct 27, 1994
727 days to decisionK925510 · Product code: **DSI** · Cardiovascular
Source: <https://www.510kdatabase.net/k925510/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Oct 30, 1992
Decision date	Oct 27, 1994
Days to decision	727 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k925510/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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