

**K864507 V-PACE(TM) TRANSLUMINAL VENTRICULAR/ MODEL
SP5540**Mar 6, 1987
112 days to decisionK864507 · Product code: **LDF** · Cardiovascular
Source: <https://www.510kdatabase.net/k864507/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	Nov 14, 1986
Decision date	Mar 6, 1987
Days to decision	112 days
Third-party review	No

APPLICANT

Company	Spectramed, Inc.
Location	Findley, MN, US
Contact	CONSTANCE G BUNDY
510(k) history	13 submissions · 13 cleared · 1987-1990

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k864507/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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