

K924390 ANGEMED 5000 LEAD ADAPTERAug 25, 1993
359 days to decisionK924390 · Product code: **LDF** · Cardiovascular
Source: <https://www.510kdatabase.net/k924390/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	Aug 31, 1992
Decision date	Aug 25, 1993
Days to decision	359 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Angemed, Inc.
Location	Minneapolis, MN, US
Contact	TOM THOMPSON
510(k) history	2 submissions · 2 cleared · 1993-1993

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