FDA 510(k) Premarket Notification

K924390 ANGEMED 5000 LEAD ADAPTER

CLEARED

Aug 25, 1993
359 days to decision

K924390 · 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