FDA 510(k) Premarket Notification

K882045 HIGH TORQUE TEMPORARY PERVENOUS LEAD W/DEPTH MARK

CLEARED

Jul 12, 1988
57 days to decision

K882045 · Product code: **LDF** · Cardiovascular
Source: https://www.510kdatabase.net/k882045/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	May 16, 1988
Decision date	Jul 12, 1988
Days to decision	57 days
Third-party review	No

APPLICANT

Company	**Cordis Leads, Inc.**
Location	Suffield, CT, US
Contact	DUANE A SCHULTZ
510(k) history	4 submissions · 4 cleared · 1988-1988