FDA 510(k) Premarket Notification

K944957 TEMPORARY MYOCARDIAL PACING LEAD, MODEL 6500

CLEARED

Jul 14, 1995
280 days to decision

K944957 · Product code: **LDF** · Cardiovascular
Source: https://www.510kdatabase.net/k944957/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	Oct 7, 1994
Decision date	Jul 14, 1995
Days to decision	280 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Medtronic Blood Systems, Inc.**
Location	Anaheim, CA, US
Contact	DIANA SALDITT
510(k) history	15 submissions · 15 cleared · 1988-1995