FDA 510(k) Premarket Notification

K922269 UNIPOLAR TEMPORARY MYOCARDIAL PACING WIRE, 6493

CLEARED

Nov 23, 1992
193 days to decision

K922269 · Product code: **LDF** · Cardiovascular
Source: https://www.510kdatabase.net/k922269/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	May 14, 1992
Decision date	Nov 23, 1992
Days to decision	193 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**The Heart Valve Div. Medtronic Cardiovascular Surg**
Location	Minneapolis, MN, US
Contact	DIANA K SALDITT
510(k) history	5 submissions · 4 cleared · 1991-1998