FDA 510(k) Premarket Notification

K954809 UNIPOLAR TEMPORARY MYOCARDIAL PACING WIRE MODEL 6494

CLEARED

Dec 13, 1995
55 days to decision

K954809 · Product code: **LDF** · Cardiovascular
Source: https://www.510kdatabase.net/k954809/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	Oct 19, 1995
Decision date	Dec 13, 1995
Days to decision	55 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Medtronic Vascular**
Location	Walker, MI, US
Contact	VALERIE K STUHR
510(k) history	475 submissions · 453 cleared · 1977-2023