

**K140972 UNIPOLAR PEDIATRIC TEMPORARY PACING LEAD,
UNIPOLAR TEMPORARY ATRIAL PACING LEAD, UNIPOLAR
TEMPORARY MYOCARDIAL PACING**Jun 6, 2014
51 days to decisionK140972 · Product code: **LDF** · Cardiovascular
Source: <https://www.510kdatabase.net/k140972/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	Apr 16, 2014
Decision date	Jun 6, 2014
Days to decision	51 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Medtronic, Inc.
Location	Mounds View, MN, US
Contact	DEBRA TAITAGUE
Website	https://www.medtronic.com
510(k) history	209 submissions · 208 cleared · 1981-2026

Medtronic, Inc. is a global medical device manufacturer headquartered in Mounds View, United States. The company develops and markets a broad range of medical devices across multiple therapeutic areas. Medtronic maintains a strong FDA 510(k) regulatory track record with FDA 510(k) cleared devices from total submissions since 1981. The company specializes primarily in Cardiovascular devices, which represent 82% of its submission portfolio. Recent clearances include coronary perfusion cannulae, intracoronary shunts, venous cannulae, guidewires, deflectable catheter systems,...