

**K862459 MODIFIED CIRCUIT BURN-IN PROCESS
IMPLANTABLE PULSE**Jan 2, 1987
186 days to decisionK862459 · Product code: **DXY** · Cardiovascular
Source: <https://www.510kdatabase.net/k862459/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Jun 30, 1986
Decision date	Jan 2, 1987
Days to decision	186 days
Third-party review	No

APPLICANT

Company	Medtronic Vascular
Location	Walker, MI, US
Contact	TIMOTHY SAMSEL
510(k) history	475 submissions · 453 cleared · 1977-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k862459/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 15, 2026