

K800069 MEDTRONIC MODELS 5966/5967, ETC.Jun 26, 1980
167 days to decisionK800069 · Product code: **DXY** · Cardiovascular
Source: <https://www.510kdatabase.net/k800069/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Jan 11, 1980
Decision date	Jun 26, 1980
Days to decision	167 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k800069/> 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