

K884215 MINIX 8340, 8341, 8342 MINIX-ST 8330, 8331Nov 17, 1988
42 days to decisionK884215 · Product code: **DXY** · CardiovascularSource: <https://www.510kdatabase.net/k884215/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Oct 6, 1988
Decision date	Nov 17, 1988
Days to decision	42 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k884215/>; 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 14, 2026