

K893633 MODEL 5330 EXTERNAL A-V SEQUENTIAL DEMAND PULSEJun 20, 1989
40 days to decisionK893633 · Product code: **DTE** · Cardiovascular
Source: <https://www.510kdatabase.net/k893633/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pulse-generator, Pacemaker, External (DTE)
Date received	May 11, 1989
Decision date	Jun 20, 1989
Days to decision	40 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k893633/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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