FDA 510(k) Premarket Notification

K893633 MODEL 5330 EXTERNAL A-V SEQUENTIAL DEMAND PULSE

CLEARED

Jun 20, 1989
40 days to decision

K893633 · 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