

K895495 ADD'L LABELING TO MINIX MODELS AND MINIX ST MODELS

Sep 26, 1989
29 days to decision

K895495 · Product code: **DXY** · Cardiovascular
Source: <https://www.510kdatabase.net/k895495/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Aug 28, 1989
Decision date	Sep 26, 1989
Days to decision	29 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k895495/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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