FDA 510(k) Premarket Notification

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

CLEARED

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