FDA 510(k) Premarket Notification

K884215 MINIX 8340, 8341, 8342 MINIX-ST 8330, 8331

CLEARED

Nov 17, 1988
42 days to decision

K884215 · Product code: **DXY** · Cardiovascular
Source: 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