FDA 510(k) Premarket Notification

K904892 VACUUM BAKE CYCLE

CLEARED

Dec 20, 1990
51 days to decision

K904892 · Product code: **DXY** · Cardiovascular
Source: https://www.510kdatabase.net/k904892/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Oct 30, 1990
Decision date	Dec 20, 1990
Days to decision	51 days
Third-party review	No

APPLICANT

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