FDA 510(k) Premarket Notification

K873158 MODIFIED MULTICOR AND OMNI-STANICOR PACEMAKERS

CLEARED

Jan 7, 1988
150 days to decision

K873158 · Product code: **DXY** · Cardiovascular
Source: https://www.510kdatabase.net/k873158/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Aug 10, 1987
Decision date	Jan 7, 1988
Days to decision	150 days
Third-party review	No

APPLICANT

Company	**Tpl-Cordis, Inc.**
Location	Suffield, CT, US
Contact	WILLIAM C NEALON
510(k) history	6 submissions · 6 cleared · 1987-1989