FDA 510(k) Premarket Notification

K870610 RESUBMISSION OF MODEL 030-437 IMPLANT. ELECTRODE

CLEARED

Mar 20, 1987
46 days to decision

K870610 · Product code: **DTB** · Cardiovascular
Source: https://www.510kdatabase.net/k870610/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Permanent Pacemaker Electrode (DTB)
Date received	Feb 2, 1987
Decision date	Mar 20, 1987
Days to decision	46 days
Third-party review	No

APPLICANT

Company	**Telectronics, Inc.**
Location	Mchenry, IL, US
Contact	WILLIAM C NEALON
510(k) history	107 submissions · 107 cleared · 1977-1990