FDA 510(k) Premarket Notification

K791727 IMPLANTABLE VENTRICULAR-INHIBITED

CLEARED

Sep 27, 1979
27 days to decision

K791727 · Product code: **DXY** · Cardiovascular
Source: https://www.510kdatabase.net/k791727/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Aug 31, 1979
Decision date	Sep 27, 1979
Days to decision	27 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k791727/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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