FDA 510(k) Premarket Notification

K831859 PACEMAKER PROGRAMMER #2032

CLEARED

Oct 27, 1983
139 days to decision

K831859 · Product code: **DXY** · Cardiovascular
Source: https://www.510kdatabase.net/k831859/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Jun 10, 1983
Decision date	Oct 27, 1983
Days to decision	139 days
Third-party review	No

APPLICANT

Company	**Vitatron Medical BV**
Location	Mchenry, IL, US
510(k) history	19 submissions · 19 cleared · 1976-1986