FDA 510(k) Premarket Notification

K880422 MAESTRO(R) MODEL 119, SINGLE-CHAMBER PACEMAKER

CLEARED

Mar 31, 1988
59 days to decision

K880422 · Product code: **DXY** · Cardiovascular
Source: https://www.510kdatabase.net/k880422/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Feb 1, 1988
Decision date	Mar 31, 1988
Days to decision	59 days
Third-party review	No

APPLICANT

Company	**Cardiac Control Systems, Inc.**
Location	Mchenry, IL, US
Contact	Scott J Mindrebo
510(k) history	26 submissions · 24 cleared · 1983-1997