

**K880326 MAESTRO(R) MODELS 115, 118, 215 AND 218
PACEMAKERS**Apr 11, 1988
76 days to decisionK880326 · Product code: **DXY** · Cardiovascular
Source: <https://www.510kdatabase.net/k880326/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Jan 26, 1988
Decision date	Apr 11, 1988
Days to decision	76 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k880326/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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