

**K880422 MAESTRO(R) MODEL 119, SINGLE-CHAMBER  
PACEMAKER**Mar 31, 1988  
59 days to decisionK880422 · Product code: **DXY** · Cardiovascular  
Source: <https://www.510kdatabase.net/k880422/>**SUBMISSION DETAILS**

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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**

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

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k880422/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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