

**K884023 MAESTRO(R) SERIES 100/200 SINGLE CHAMBER
PACEMAKER**Oct 17, 1988
25 days to decisionK884023 · Product code: **DXY** · Cardiovascular
Source: <https://www.510kdatabase.net/k884023/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Sep 22, 1988
Decision date	Oct 17, 1988
Days to decision	25 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k884023/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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