

K904941 MAESTRO SERIES 100/200 CARDIAC SINGLE CHAMBER PACEJan 3, 1991
63 days to decisionK904941 · Product code: **DXY** · Cardiovascular
Source: <https://www.510kdatabase.net/k904941/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Nov 1, 1990
Decision date	Jan 3, 1991
Days to decision	63 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/k904941/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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