

K770716 GENERATOR, PULSE, UNIPOLAR, MODEL #504Apr 26, 1977
8 days to decisionK770716 · Product code: **DXY** · CardiovascularSource: <https://www.510kdatabase.net/k770716/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Apr 18, 1977
Decision date	Apr 26, 1977
Days to decision	8 days
Third-party review	No

APPLICANT

Company	Cardiac Pacemakers, Inc.
Location	Mchenry, IL, US
510(k) history	76 submissions · 76 cleared · 1977-2010

Cardiac Pacemakers, Inc. (CPI), doing business as Guidant Cardiac Rhythm Management, manufactured implantable cardiac rhythm management devices. Now part of Boston Scientific, the company is based in Saint Paul, Minnesota, with historical operations in McHenry, US. The company received FDA 510(k) clearances from total submissions between 1977 and 2010. Cardiovascular devices dominated the regulatory portfolio at 83% of submissions. This historical record reflects the company's core focus on cardiac rhythm management and related interventional technologies. CPI developed t...

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Device record: <https://www.510kdatabase.net/k770716/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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