

**K771992 L-501 CARDIAC PACEMAKER**Nov 2, 1977  
12 days to decisionK771992 · Product code: **DXY** · CardiovascularSource: <https://www.510kdatabase.net/k771992/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Oct 21, 1977
Decision date	Nov 2, 1977
Days to decision	12 days
Third-party review	No

**APPLICANT**

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Company	<b>Coratomic, Inc.</b>
Location	Mchenry, IL, US
510(k) history	15 submissions · 14 cleared · 1976-1986

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Device record: <https://www.510kdatabase.net/k771992/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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