

**K791762 SELF CHECK**Sep 21, 1979  
24 days to decisionK791762 · Product code: **DTC** · CardiovascularSource: <https://www.510kdatabase.net/k791762/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Analyzer, Pacemaker Generator Function (DTC)
Date received	Aug 28, 1979
Decision date	Sep 21, 1979
Days to decision	24 days
Third-party review	No

**APPLICANT**

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Company	<b>Instromedix, Inc.</b>
Location	Mchenry, IL, US
510(k) history	32 submissions · 32 cleared · 1977-1997

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k791762/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 29, 2026