

**K781111 LEAD CONVERSION KIT, MODEL 341**Aug 31, 1978  
59 days to decision

K781111 · Cardiovascular

Source: <https://www.510kdatabase.net/k781111/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Date received	Jul 3, 1978
Decision date	Aug 31, 1978
Days to decision	59 days
Third-party review	No

**APPLICANT**

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Company	<b>Edwards Pacemaker Systems</b>
Location	Mchenry, IL, US
510(k) history	13 submissions · 13 cleared · 1977-1979

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k781111/> 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 July 9, 2026