

**K874204 450SLL DEFIBRILLATOR**Jan 28, 1988  
105 days to decisionK874204 · Product code: **LDD** · CardiovascularSource: <https://www.510kdatabase.net/k874204/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Oct 15, 1987
Decision date	Jan 28, 1988
Days to decision	105 days
Third-party review	No

**APPLICANT**

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Company	<b>Medical Research Laboratories, Inc.</b>
Location	Mchenry, IL, US
Contact	DILIP MEHTA
510(k) history	19 submissions · 15 cleared · 1981-2002

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k874204/>; 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 May 25, 2026