

**K904466 MODEL #250 DEFIB CABLE**Jan 30, 1991  
121 days to decisionK904466 · Product code: **LDD** · Cardiovascular  
Source: <https://www.510kdatabase.net/k904466/>**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 1, 1990
Decision date	Jan 30, 1991
Days to decision	121 days
Third-party review	No

**APPLICANT**

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Company	<b>Cardiotronics, Inc.</b>
Location	West Carlsbad, CA, US
Contact	TIM J WAY
510(k) history	27 submissions · 27 cleared · 1988-1995

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k904466/>; 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 6, 2026