

**K910634 DC200 DEFIBRILLATOR**Jul 24, 1991  
161 days to decisionK910634 · Product code: **LDD** · CardiovascularSource: <https://www.510kdatabase.net/k910634/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Feb 13, 1991
Decision date	Jul 24, 1991
Days to decision	161 days
Third-party review	No

**APPLICANT**

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Company	<b>Siemens Burdick, Inc.</b>
Location	Miltoon, WI, US
Contact	PAUL E APPEL
510(k) history	7 submissions · 7 cleared · 1991-1992

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k910634/>; 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 30, 2026