

**K863303 MODEL 617D**Sep 5, 1986  
11 days to decisionK863303 · Product code: **LDD** · CardiovascularSource: <https://www.510kdatabase.net/k863303/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Aug 25, 1986
Decision date	Sep 5, 1986
Days to decision	11 days
Third-party review	No

**APPLICANT**

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Company	<b>Quinton, Inc.</b>
Location	Mchenry, IL, US
Contact	RON R DUCK
510(k) history	164 submissions · 160 cleared · 1976-2003

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k863303/>; 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 20, 2026