

**K895185 FIRST MEDIC MODEL 610P**Oct 27, 1989  
77 days to decisionK895185 · Product code: **LDD** · CardiovascularSource: <https://www.510kdatabase.net/k895185/>**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 11, 1989
Decision date	Oct 27, 1989
Days to decision	77 days
Third-party review	No

**APPLICANT**

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Company	<b>First Medical Devices Corp.</b>
Location	Bellevue, WA, US
Contact	BRUCE G HAGGAR
510(k) history	7 submissions · 7 cleared · 1986-1990

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k895185/>; 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