

**K903018 MEDTRONIC ANDOVER MEDICAL SAF-D-FIB MODEL  
1220-002**Dec 3, 1990  
145 days to decisionK903018 · Product code: **LDD** · Cardiovascular  
Source: <https://www.510kdatabase.net/k903018/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Jul 11, 1990
Decision date	Dec 3, 1990
Days to decision	145 days
Third-party review	No

**APPLICANT**

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Company	<b>Medtronic Andover Medical, Inc.</b>
Location	Lowell, MA, US
Contact	JANICE PEVIDE
510(k) history	6 submissions · 6 cleared · 1985-1991

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k903018/>; 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 14, 2026