

K903657 MODEL 610 MODIFIEDSep 4, 1990
26 days to decisionK903657 · Product code: **LDD** · CardiovascularSource: <https://www.510kdatabase.net/k903657/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Aug 9, 1990
Decision date	Sep 4, 1990
Days to decision	26 days
Third-party review	No

APPLICANT

Company	First Medical Devices Corp.
Location	Bellevue, WA, US
Contact	RAYMOND W GIFFORD
510(k) history	7 submissions · 7 cleared · 1986-1990

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