

K882235 POLY-TACOct 5, 1988
131 days to decisionK882235 · Product code: **LDD** · Cardiovascular
Source: <https://www.510kdatabase.net/k882235/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	May 27, 1988
Decision date	Oct 5, 1988
Days to decision	131 days
Third-party review	No

APPLICANT

Company	Cardiovascular Group, Inc.
Location	Lake Oswego, OR, US
Contact	JOSEPH M MERTZ
510(k) history	1 submissions · 1 cleared · 1988-1988

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