

K853354 MEDIC 4Oct 11, 1985
60 days to decisionK853354 · Product code: **LDD** · CardiovascularSource: <https://www.510kdatabase.net/k853354/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Aug 12, 1985
Decision date	Oct 11, 1985
Days to decision	60 days
Third-party review	No

APPLICANT

Company	Kone Instruments, Inc.
Location	Mchenry, IL, US
Contact	FRANCIS I DOMINY
510(k) history	12 submissions · 12 cleared · 1982-1987

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