

K843229 LIFEPAK 200Oct 18, 1984
62 days to decisionK843229 · Product code: **LDD** · CardiovascularSource: <https://www.510kdatabase.net/k843229/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Aug 17, 1984
Decision date	Oct 18, 1984
Days to decision	62 days
Third-party review	No

APPLICANT

Company	Physio-Control Corp.
Location	Mchenry, IL, US
Contact	MICHAEL B GROPP
510(k) history	80 submissions · 78 cleared · 1976-1999

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