

K890079 LIFEPAK 10 DEFIBRILLATOR/MONITORFeb 23, 1989
50 days to decisionK890079 · Product code: **LDD** · Cardiovascular
Source: <https://www.510kdatabase.net/k890079/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Jan 4, 1989
Decision date	Feb 23, 1989
Days to decision	50 days
Third-party review	No

APPLICANT

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

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