

**K931053 LIFEPAK 10 DEFIBRILLATOR/MONITOR W/OPT
PACEMAKER**May 18, 1993
71 days to decisionK931053 · Product code: **LDD** · Cardiovascular
Source: <https://www.510kdatabase.net/k931053/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Mar 8, 1993
Decision date	May 18, 1993
Days to decision	71 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k931053/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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