

K902288 MODIFIED LIFEPAK 9 MONITOR/DEFIBRILLATORFeb 25, 1991
279 days to decisionK902288 · Product code: **LDD** · CardiovascularSource: <https://www.510kdatabase.net/k902288/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	May 22, 1990
Decision date	Feb 25, 1991
Days to decision	279 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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