

K920990 LIFE-TEL L.B.E.(LIFE-PAK 5/10 BATTERY ELIMINATOR)Jul 6, 1992
126 days to decisionK920990 · Product code: **LDD** · Cardiovascular
Source: <https://www.510kdatabase.net/k920990/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Mar 2, 1992
Decision date	Jul 6, 1992
Days to decision	126 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Raco Products Co.
Location	Cleveland, OH, US
Contact	GERALD J CAHILL
510(k) history	4 submissions · 4 cleared · 1992-1994

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k920990/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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