

**K940842 UNIPOWER RECHARGEABLE BATTERY PACK**Sep 7, 1994  
196 days to decisionK940842 · Product code: **LDD** · Cardiovascular  
Source: <https://www.510kdatabase.net/k940842/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - ST
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Feb 23, 1994
Decision date	Sep 7, 1994
Days to decision	196 days
Third-party review	No

**APPLICANT**

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Company	<b>Unipower Corp.</b>
Location	Minneapolis, MN, US
Contact	L.E PERTL
510(k) history	3 submissions · 2 cleared · 1994-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k940842/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 26, 2026