

**K832833 PERSONAL DEFIBRILLATOR**Oct 26, 1984  
434 days to decisionK832833 · Product code: **LDD** · CardiovascularSource: <https://www.510kdatabase.net/k832833/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Aug 19, 1983
Decision date	Oct 26, 1984
Days to decision	434 days
Third-party review	No

**APPLICANT**

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Company	<b>Physio-Control, Inc.</b>
Location	Redmond, WA, US
Contact	SYLVIA LEMKE
510(k) history	14 submissions · 14 cleared · 1984-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k832833/>; 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