

**K822617 VPD540 DEFIBRILLATOR**Apr 28, 1983  
241 days to decisionK822617 · Product code: **LDD** · CardiovascularSource: <https://www.510kdatabase.net/k822617/>**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 30, 1982
Decision date	Apr 28, 1983
Days to decision	241 days
Third-party review	No

**APPLICANT**

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Company	<b>Life Science Instrumentation, Inc.</b>
Location	Mchenry, IL, US
510(k) history	20 submissions · 20 cleared · 1981-1985

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k822617/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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