

**K930548 DPD DEFIBRILLATOR**Jun 25, 1993  
142 days to decisionK930548 · Product code: **LDD** · Cardiovascular  
Source: <https://www.510kdatabase.net/k930548/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Feb 3, 1993
Decision date	Jun 25, 1993
Days to decision	142 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>Medical Research Laboratories, Inc.</b>
Location	Mchenry, IL, US
Contact	DILIP MEHTA
510(k) history	19 submissions · 15 cleared · 1981-2002

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k930548/> 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 25, 2026