

**K874091 360D DC DEFIBRILLATOR**Nov 23, 1987  
47 days to decisionK874091 · Product code: **LDD** · Cardiovascular  
Source: <https://www.510kdatabase.net/k874091/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Oct 7, 1987
Decision date	Nov 23, 1987
Days to decision	47 days
Third-party review	No

**APPLICANT**

---

Company	<b>Medac, Inc.</b>
Location	Tualatin, OR, US
Contact	ROBERT M BOONSTRA
510(k) history	10 submissions · 10 cleared · 1987-1988

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k874091/>, 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 July 7, 2026