

**K081574 ZOLL R SERIES**Sep 18, 2008  
105 days to decisionK081574 · Product code: **LDD** · Cardiovascular  
Source: <https://www.510kdatabase.net/k081574/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Jun 5, 2008
Decision date	Sep 18, 2008
Days to decision	105 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>ZOLL Medical Corporation</b>
Location	Chelmsford, MA, US
Contact	EILEEN M BOYLE
510(k) history	30 submissions · 30 cleared · 2005-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k081574/> 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 June 6, 2026