

**K100654 ZOLL PROPAQ MD**Jul 29, 2010  
143 days to decisionK100654 · Product code: **LDD** · Cardiovascular  
Source: <https://www.510kdatabase.net/k100654/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Mar 8, 2010
Decision date	Jul 29, 2010
Days to decision	143 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Zoll Medical Corporation, World Wide Headquarters</b>
Location	Chelmsford, MA, US
Contact	PAUL DIAS
510(k) history	21 submissions · 21 cleared · 2007-2015

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k100654/> 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 5, 2026