

**K063602 AUTOPULSE RESUSCITATION SYSTEM MODEL100**Dec 21, 2006  
17 days to decisionK063602 · Product code: **DRM** · Cardiovascular  
Source: <https://www.510kdatabase.net/k063602/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Compressor, Cardiac, External (DRM)
Date received	Dec 4, 2006
Decision date	Dec 21, 2006
Days to decision	17 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Zoll Circulation</b>
Location	Sunnyvale, CA, US
Contact	MARK PERKINS
510(k) history	5 submissions · 4 cleared · 2006-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k063602/> 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 13, 2026