

**K040453 AUTOPULSE RESUSCITATION SYSTEM, MODEL 100**Mar 11, 2004  
17 days to decisionK040453 · Product code: **DRM** · Cardiovascular  
Source: <https://www.510kdatabase.net/k040453/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Compressor, Cardiac, External (DRM)
Date received	Feb 23, 2004
Decision date	Mar 11, 2004
Days to decision	17 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Revivant Corp.</b>
Location	Sunnyvale, CA, US
Contact	BOB H KATZ
510(k) history	4 submissions · 4 cleared · 2001-2004

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k040453/> 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 3, 2026