

**K062977 CRITIVIEW, MODEL CRV3**Feb 28, 2007  
152 days to decisionK062977 · Product code: **DPW** · CardiovascularSource: <https://www.510kdatabase.net/k062977/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Flowmeter, Blood, Cardiovascular (DPW)
Date received	Sep 29, 2006
Decision date	Feb 28, 2007
Days to decision	152 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Critisense , Ltd.</b>
Location	Mccordsville, IN, US
Contact	PAUL DRYDEN
510(k) history	2 submissions · 2 cleared · 2006-2007

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k062977/> 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 28, 2026