

**K041180 QUANTIX OR VESSEL STABILIZER DEVICE**Jun 7, 2004  
32 days to decisionK041180 · Product code: **DPW** · CardiovascularSource: <https://www.510kdatabase.net/k041180/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Flowmeter, Blood, Cardiovascular (DPW)
Date received	May 6, 2004
Decision date	Jun 7, 2004
Days to decision	32 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Cardiosonix, Ltd.</b>
Location	Kfar Saba, IL
Contact	AHAVA STEIN
510(k) history	3 submissions · 3 cleared · 2002-2004

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k041180/> 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