

**K903633 ADVANCE LASER FLOWMETER, ALF 21**Oct 9, 1990  
60 days to decisionK903633 · Product code: **DPW** · CardiovascularSource: <https://www.510kdatabase.net/k903633/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Flowmeter, Blood, Cardiovascular (DPW)
Date received	Aug 10, 1990
Decision date	Oct 9, 1990
Days to decision	60 days
Third-party review	No

**APPLICANT**

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Company	<b>Transonic Systems, Inc.</b>
Location	Ithaca, NY, US
Contact	CORNELIS J DROST
510(k) history	16 submissions · 16 cleared · 1988-2015

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k903633/>; 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 4, 2026