

**K904317 AVALON GENERAL PURPOSE SCANNER, PREVIEW  
GP**Feb 13, 1991  
146 days to decisionK904317 · Product code: **DPW** · Cardiovascular  
Source: <https://www.510kdatabase.net/k904317/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Flowmeter, Blood, Cardiovascular (DPW)
Date received	Sep 20, 1990
Decision date	Feb 13, 1991
Days to decision	146 days
Third-party review	No

**APPLICANT**

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Company	<b>Avalon Technology Corp.</b>
Location	Indianapolis, IN, US
Contact	WILLIAM D LIPPS
510(k) history	2 submissions · 2 cleared · 1989-1991

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k904317/>; 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 May 21, 2026