

**K926219 516YT DISPLAY DRIVER**Dec 16, 1993  
371 days to decisionK926219 · Product code: **DXJ** · Cardiovascular  
Source: <https://www.510kdatabase.net/k926219/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Display, Cathode-ray Tube, Medical (DXJ)
Date received	Dec 10, 1992
Decision date	Dec 16, 1993
Days to decision	371 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Vidco, Inc.</b>
Location	Beaverton, OR, US
Contact	BRADLEY A COHEN
510(k) history	7 submissions · 7 cleared · 1989-1997

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k926219/> 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 4, 2026