

**K933517 AUTOMATIC PULL BACK DEVICE**Oct 15, 1993  
87 days to decisionK933517 · Product code: **DQX** · CardiovascularSource: <https://www.510kdatabase.net/k933517/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Jul 20, 1993
Decision date	Oct 15, 1993
Days to decision	87 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Cardiovascular Imaging Systems, Inc.</b>
Location	Sunnyvale, CA, US
Contact	YUE-TEH JANG
510(k) history	19 submissions · 19 cleared · 1989-1997

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k933517/> 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 20, 2026