

**K930961 CATHETER GUIDEWIRE (PERIPHERAL USE)**May 26, 1993  
91 days to decisionK930961 · Product code: **DQX** · Cardiovascular  
Source: <https://www.510kdatabase.net/k930961/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Feb 24, 1993
Decision date	May 26, 1993
Days to decision	91 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Target Therapeutics</b>
Location	Los Angeles, CA, US
Contact	GRACE CARLAND
510(k) history	70 submissions · 70 cleared · 1985-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k930961/> 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 30, 2026