

**K030617 PT2 GUIDE WIRE AND ADDWIRE EXTENSION WIRE**May 21, 2003  
84 days to decisionK030617 · Product code: **DQX** · CardiovascularSource: <https://www.510kdatabase.net/k030617/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Wire, Guide, Catheter (DQX)
Date received	Feb 26, 2003
Decision date	May 21, 2003
Days to decision	84 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Boston Scientific Scimed, Inc.</b>
Location	Plymouth, MN, US
Contact	ANNE V ROSSI
510(k) history	35 submissions · 26 cleared · 1994-2004

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k030617/> 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 28, 2026