

**K961999 SCHNEIDER GUIDER SOFTIP GUIDING CATHETERS**Nov 19, 1996  
181 days to decisionK961999 · Product code: **DYG** · CardiovascularSource: <https://www.510kdatabase.net/k961999/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Flow Directed (DYG)
Date received	May 22, 1996
Decision date	Nov 19, 1996
Days to decision	181 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	MARIA BRITTLE
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/k961999/> 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