

**K924591 SCHNEIDER SOLID-7 SOPTIP GUIDING CATHETER**Dec 8, 1992  
89 days to decisionK924591 · Product code: **DQY** · CardiovascularSource: <https://www.510kdatabase.net/k924591/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Sep 10, 1992
Decision date	Dec 8, 1992
Days to decision	89 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Schneider Intl., Ltd.</b>
Location	Minneapolis, MN, US
Contact	ROBERT L ULLEN
510(k) history	22 submissions · 22 cleared · 1989-1995

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k924591/> 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 1, 2026