

**K915347 SCIMED VIPER CATHETER**Aug 17, 1992  
265 days to decisionK915347 · Product code: LIT · Cardiovascular  
Source: <https://www.510kdatabase.net/k915347/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Angioplasty, Peripheral, Transluminal (LIT)
Date received	Nov 26, 1991
Decision date	Aug 17, 1992
Days to decision	265 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Scimed</b>
Location	Maple Grove, MN, US
Contact	DARLENE THOMETZ
510(k) history	8 submissions · 8 cleared · 1990-1999

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k915347/> 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 4, 2026