

**K931421 DISPOSABLE STEERABLE ANGIOSCOPE**Jun 16, 1993  
86 days to decisionK931421 · Product code: **LYK** · CardiovascularSource: <https://www.510kdatabase.net/k931421/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Angioscope (LYK)
Date received	Mar 22, 1993
Decision date	Jun 16, 1993
Days to decision	86 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Endovascular, Inc.</b>
Location	Costa Mesa, CA, US
Contact	KAREN U SALINAS
510(k) history	3 submissions · 3 cleared · 1993-1993

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k931421/> 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 6, 2026