

**K926293 ANGIOCOR SIDEHOLE INFUSION CATHETER**Mar 7, 1994  
448 days to decisionK926293 · Product code: **LJS** · General Hospital  
Source: <https://www.510kdatabase.net/k926293/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Long-term Greater Than 30 Days (LJS)
Date received	Dec 14, 1992
Decision date	Mar 7, 1994
Days to decision	448 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Angiocor</b>
Location	Great Neck, NY, US
Contact	ANAND AKERKAR
510(k) history	1 submissions · 1 cleared · 1994-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k926293/> 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