

**K010721 AXIOM ARTIS MODULAR ANGIOGRAPHY SYSTEM**Mar 30, 2001  
18 days to decisionK010721 · Product code: **IZI** · RadiologySource: <https://www.510kdatabase.net/k010721/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Angiographic (IZI)
Date received	Mar 12, 2001
Decision date	Mar 30, 2001
Days to decision	18 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Siemens Medical Corp.</b>
Location	Iselin, NJ, US
Contact	PRAVEEN NADKARNI
510(k) history	5 submissions · 5 cleared · 1999-2001

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k010721/> 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 May 19, 2026