

**K052039 GE OEC ALTITUDE**Aug 11, 2005  
14 days to decisionK052039 · Product code: **IZI** · Radiology  
Source: <https://www.510kdatabase.net/k052039/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	System, X-ray, Angiographic (IZI)
Date received	Jul 28, 2005
Decision date	Aug 11, 2005
Days to decision	14 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Ge Oec Medical Systems</b>
Location	Salt Lake City, UT, US
Contact	SUSAN SCHMIDT
510(k) history	7 submissions · 7 cleared · 2001-2008

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k052039/> 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 18, 2026