

**K960575 ADVANTX LCV+**May 7, 1996  
85 days to decisionK960575 · Product code: **IZI** · Radiology  
Source: <https://www.510kdatabase.net/k960575/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Angiographic (IZI)
Date received	Feb 12, 1996
Decision date	May 7, 1996
Days to decision	85 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>GE Medical Systems</b>
Location	Milwaukee, WI, US
Contact	LARRY A KROGER, PH.D.
510(k) history	169 submissions · 166 cleared · 1989-2022

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k960575/> 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 27, 2026