

**K070089 SYNTERMED LIVE**Mar 2, 2007  
52 days to decisionK070089 · Product code: **LLZ** · Radiology  
Source: <https://www.510kdatabase.net/k070089/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Jan 9, 2007
Decision date	Mar 2, 2007
Days to decision	52 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Syntermed, Inc.</b>
Location	Atlanta, GA, US
Contact	KENNETH F VAN TRAIN
510(k) history	12 submissions · 12 cleared · 2000-2013

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k070089/> 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 17, 2026