

**K161419 Multi Modality Viewer**Jul 13, 2016  
51 days to decisionK161419 · Product code: **LLZ** · Radiology  
Source: <https://www.510kdatabase.net/k161419/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	May 23, 2016
Decision date	Jul 13, 2016
Days to decision	51 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Vital Images, Inc.</b>
Location	St. Paul, MN, US
Contact	KATIE RYAN
510(k) history	41 submissions · 41 cleared · 1995-2018

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k161419/> 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 June 18, 2026