

**K071492 FIRSTVIEW**Aug 28, 2007  
90 days to decisionK071492 · Product code: **LLZ** · Radiology  
Source: <https://www.510kdatabase.net/k071492/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	May 30, 2007
Decision date	Aug 28, 2007
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Riverain Medical Group,Llc</b>
Location	Miamisburg, OH, US
Contact	JENNIFER STEINKE
510(k) history	2 submissions · 2 cleared · 2007-2011

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k071492/> 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