

**K092363 SOFTVIEW, MODEL 2.01**Mar 18, 2010  
225 days to decisionK092363 · Product code: **LLZ** · Radiology  
Source: <https://www.510kdatabase.net/k092363/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Aug 5, 2009
Decision date	Mar 18, 2010
Days to decision	225 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Riverain Medical Group</b>
Location	Miamisburg, OH, US
Contact	JENNIFER VETTER
510(k) history	1 submissions · 1 cleared · 2010-2010

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k092363/> 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