

**K101842 ACIES**Feb 4, 2011  
218 days to decisionK101842 · Product code: **LLZ** · Radiology  
Source: <https://www.510kdatabase.net/k101842/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Jul 1, 2010
Decision date	Feb 4, 2011
Days to decision	218 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Konica Minolta Medical &amp; Graphic, Inc.</b>
Location	Hachioji-Shi Tokyo, JP
Contact	RUSSELL MUNVES
510(k) history	27 submissions · 27 cleared · 2002-2013

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k101842/> 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