

**K091602 MUCHECK V8**Jun 18, 2009  
15 days to decisionK091602 · Product code: **IYE** · Radiology  
Source: <https://www.510kdatabase.net/k091602/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accelerator, Linear, Medical (IYE)
Date received	Jun 3, 2009
Decision date	Jun 18, 2009
Days to decision	15 days
Third-party review	Yes
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Oncology Data Systems, Inc.</b>
Location	Oklahoma City, OK, US
Contact	VINCE RUMINER
510(k) history	4 submissions · 4 cleared · 2001-2010

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k091602/> 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