

**K111862 VIEWRAY SYSTEM**May 22, 2012  
327 days to decisionK111862 · Product code: **IYE** · Radiology  
Source: <https://www.510kdatabase.net/k111862/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accelerator, Linear, Medical (IYE)
Date received	Jun 30, 2011
Decision date	May 22, 2012
Days to decision	327 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Viewray, Incorporated</b>
Location	Oakwood Village, OH, US
Contact	JANICE BROWNLEE
510(k) history	6 submissions · 6 cleared · 2011-2021

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k111862/> 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 26, 2026