

**K020294 MOTION TRACKING**Apr 22, 2002  
84 days to decisionK020294 · Product code: **IYE** · Radiology  
Source: <https://www.510kdatabase.net/k020294/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accelerator, Linear, Medical (IYE)
Date received	Jan 28, 2002
Decision date	Apr 22, 2002
Days to decision	84 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Accuray, Inc.</b>
Location	Sunnyvale, CA, US
Contact	DONALD E CADDES
510(k) history	10 submissions · 10 cleared · 1999-2009

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k020294/> 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