

**K101220 EXPLORER 4D TREATMENT PLANNING SYSTEM**Nov 5, 2010  
186 days to decisionK101220 · Product code: **MUJ** · Radiology  
Source: <https://www.510kdatabase.net/k101220/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Planning, Radiation Therapy Treatment (MUJ)
Date received	May 3, 2010
Decision date	Nov 5, 2010
Days to decision	186 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>American Radiosurgery, Inc.</b>
Location	San Diego, CA, US
Contact	DAN OLIVER
510(k) history	3 submissions · 3 cleared · 2006-2010

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k101220/> 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