

**K032456 PROWESS**Oct 16, 2003  
66 days to decisionK032456 · Product code: **MUJ** · Radiology  
Source: <https://www.510kdatabase.net/k032456/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Planning, Radiation Therapy Treatment (MUJ)
Date received	Aug 11, 2003
Decision date	Oct 16, 2003
Days to decision	66 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Prowess, Inc.</b>
Location	Chico, CA, US
Contact	W. JAMES BISHOP
510(k) history	9 submissions · 9 cleared · 2003-2023

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k032456/> 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