

**K122616 PANTHER OIS/R & V MODEL VERSION 2.0**Dec 21, 2012  
116 days to decisionK122616 · Product code: **IYE** · Radiology  
Source: <https://www.510kdatabase.net/k122616/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accelerator, Linear, Medical (IYE)
Date received	Aug 27, 2012
Decision date	Dec 21, 2012
Days to decision	116 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Prowess, Inc.</b>
Location	Chico, CA, US
Contact	RACHEL SCARANO
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/k122616/> 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