

**K123746 AMES THERAPY DEVICE**May 24, 2013  
169 days to decisionK123746 · Product code: **IKK** · Physical MedicineSource: <https://www.510kdatabase.net/k123746/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Isokinetic Testing And Evaluation (IKK)
Date received	Dec 6, 2012
Decision date	May 24, 2013
Days to decision	169 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Ames Technology, Inc.</b>
Location	Eugene, OR, US
Contact	SHEILA RAMERMAN, RAC
510(k) history	1 submissions · 1 cleared · 2013-2013

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k123746/> 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 June 8, 2026