

**K072519 ACTIVATOR V SPINAL ADJUSTING INSTRUMENT**Oct 23, 2007  
46 days to decisionK072519 · Product code: **LXM** · Physical MedicineSource: <https://www.510kdatabase.net/k072519/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Manipulator, Plunger-like Joint (LXM)
Date received	Sep 7, 2007
Decision date	Oct 23, 2007
Days to decision	46 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Activator Methods International, Ltd.</b>
Location	Phoenix, AZ, US
Contact	DEBBIE KOENEMAN
510(k) history	3 submissions · 3 cleared · 2001-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k072519/> 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 15, 2026