

**K112606 ACTIVATOR V-E**Mar 6, 2012  
180 days to decisionK112606 · Product code: **LXM** · Physical Medicine  
Source: <https://www.510kdatabase.net/k112606/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Manipulator, Plunger-like Joint (LXM)
Date received	Sep 8, 2011
Decision date	Mar 6, 2012
Days to decision	180 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Activator Methods International, Ltd.</b>
Location	Phoenix, AZ, US
Contact	ARLAN W FUHR
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/k112606/> 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