

**K091000 KNEEKG**Oct 26, 2009  
201 days to decisionK091000 · Product code: **LXJ** · Physical Medicine  
Source: <https://www.510kdatabase.net/k091000/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Interactive Rehabilitation Exercise Devices (LXJ)
Date received	Apr 8, 2009
Decision date	Oct 26, 2009
Days to decision	201 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Emovi</b>
Location	Montreal, Quebec, CA
Contact	EMMANUEL MONTINI
510(k) history	1 submissions · 1 cleared · 2009-2009

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k091000/> 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 22, 2026