

**K152501 Spinal Jaxx Interbody Fusion Device**Jun 29, 2016  
302 days to decisionK152501 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k152501/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Sep 1, 2015
Decision date	Jun 29, 2016
Days to decision	302 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Neuropro Spinaljaxx, Inc.</b>
Location	Modesto, CA, US
Contact	BENJAMIN J. REMINGTON
510(k) history	1 submissions · 1 cleared · 2016-2016

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k152501/> 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 17, 2026