

**K112359 PROW FUSION**Nov 8, 2011  
84 days to decisionK112359 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k112359/>**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	Aug 16, 2011
Decision date	Nov 8, 2011
Days to decision	84 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Nlt Spine, Ltd.</b>
Location	Washington, Dc, DC, US
Contact	JOHN J SMITH
510(k) history	8 submissions · 8 cleared · 2011-2016

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k112359/> 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