

**K151654 PROW FUSION**Jul 17, 2015  
29 days to decisionK151654 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k151654/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Jun 18, 2015
Decision date	Jul 17, 2015
Days to decision	29 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	Eti Zinger
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/k151654/> 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