

**K142594 nva, nvp, nvt**Nov 28, 2014  
74 days to decisionK142594 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k142594/>**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 15, 2014
Decision date	Nov 28, 2014
Days to decision	74 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Nvision Biomedical Technologies, LLC</b>
Location	San Antonio, TN, US
Contact	Brian Kieser
510(k) history	10 submissions · 10 cleared · 2014-2019

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k142594/> 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 28, 2026