

**K121733 INFILL INTERVERTEBRAL BODY FUSION DEVICE**Jul 13, 2012  
30 days to decisionK121733 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k121733/>**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 13, 2012
Decision date	Jul 13, 2012
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Pinnacle Spine Group, LLC</b>
Location	Dallas, TX, US
Contact	Rebecca K Pine
510(k) history	12 submissions · 12 cleared · 2011-2017

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k121733/> 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 16, 2026