

**K110933 SPINEOLOGY PEEK LUMBAR INTERBODY FUSION  
DEVICE**Jun 13, 2011  
70 days to decisionK110933 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k110933/>**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	Apr 4, 2011
Decision date	Jun 13, 2011
Days to decision	70 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Spineology, Inc.</b>
Location	Stillwater, MN, US
Contact	BRYAN BECKER
510(k) history	54 submissions · 51 cleared · 1999-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k110933/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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