

**K092815 S-LIF INTERVERTEBRAL BODY FUSION DEVICE**Jun 30, 2010  
289 days to decisionK092815 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k092815/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Sep 14, 2009
Decision date	Jun 30, 2010
Days to decision	289 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Spinefrontier, Inc.</b>
Location	Beverly, MA, US
Contact	JOHN SULLIVAN
510(k) history	24 submissions · 24 cleared · 2007-2020

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k092815/> 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 15, 2026