

**K090566 PATHWAY AVID**Jun 1, 2009  
90 days to decisionK090566 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k090566/>**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	Mar 3, 2009
Decision date	Jun 1, 2009
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Custom Spine, Inc.</b>
Location	Conway, NH, US
Contact	SAAD ATTIYAH
510(k) history	12 submissions · 12 cleared · 2005-2015

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k090566/> 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