

**K090376 SPINESMITH CYNCH SPINAL SYSTEM**Apr 1, 2009  
43 days to decisionK090376 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k090376/>**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	Feb 17, 2009
Decision date	Apr 1, 2009
Days to decision	43 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Spine Smith Partners L.P.</b>
Location	Austin, TX, US
Contact	ROBERT JONES
510(k) history	9 submissions · 9 cleared · 2008-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k090376/> 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 18, 2026