

**K101085 SPINESMITH CYNCH SPINAL SYSTEM**Jul 1, 2010  
73 days to decisionK101085 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k101085/>**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	Apr 19, 2010
Decision date	Jul 1, 2010
Days to decision	73 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	LAURA LEBOEUF
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/k101085/> 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