

**K123752 ACCULIF TL AND PL CAGE**Jan 14, 2013  
39 days to decisionK123752 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k123752/>**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	Dec 6, 2012
Decision date	Jan 14, 2013
Days to decision	39 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Coalign Innovations, Inc.</b>
Location	Washington, DC, US
Contact	JUSTIN EGGLETON
510(k) history	9 submissions · 9 cleared · 2011-2013

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k123752/> 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