

**K113465 ACCULIF TL CAGE**Dec 12, 2011  
20 days to decisionK113465 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k113465/>**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	Nov 22, 2011
Decision date	Dec 12, 2011
Days to decision	20 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/k113465/> 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