

**K121683 ACCULIF TL CAGE**Jul 5, 2012  
28 days to decisionK121683 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k121683/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Jun 7, 2012
Decision date	Jul 5, 2012
Days to decision	28 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k121683/> 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