

**K140303 ALAMO T**Apr 21, 2014  
74 days to decisionK140303 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k140303/>**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	Feb 6, 2014
Decision date	Apr 21, 2014
Days to decision	74 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Alliance Partners, LLC</b>
Location	Avon Lake, OH, US
Contact	KELLEN HILLS
510(k) history	10 submissions · 10 cleared · 2011-2020

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k140303/> 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 June 18, 2026