

**K182406 Altus Spine Interbody Fusion System**Feb 22, 2019  
171 days to decisionK182406 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k182406/>**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	Sep 4, 2018
Decision date	Feb 22, 2019
Days to decision	171 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Altus Partners, LLC</b>
Location	Newtown Square, PA, US
Contact	Mark Melton
510(k) history	17 submissions · 17 cleared · 2015-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k182406/> 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 26, 2026