

**K160976 Altus Spine Interbody Fusion System**Jan 25, 2017  
293 days to decisionK160976 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k160976/>**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	Apr 7, 2016
Decision date	Jan 25, 2017
Days to decision	293 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	Claudia Hill
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/k160976/> 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