

**K162879 Elite Expandable Interbody Fusion Device**Jan 30, 2017  
108 days to decisionK162879 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k162879/>**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	Oct 14, 2016
Decision date	Jan 30, 2017
Days to decision	108 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Spineology, Inc.</b>
Location	Stillwater, MN, US
Contact	Jacqueline A. Hauge
510(k) history	54 submissions · 51 cleared · 1999-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k162879/> 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 16, 2026