

**K141942 MIDLINE II TI**Nov 7, 2014  
113 days to decisionK141942 · Product code: **OVD** · Orthopedic  
Source: <https://www.510kdatabase.net/k141942/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Integrated Fixation, Lumbar (OVD)
Date received	Jul 17, 2014
Decision date	Nov 7, 2014
Days to decision	113 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Centinel Spine, Inc.</b>
Location	West Chester, PA, US
Contact	JUSTIN EGGLETON
510(k) history	10 submissions · 10 cleared · 2012-2018

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k141942/> 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