

**K082309 CAMBRIA**Apr 9, 2009  
239 days to decisionK082309 · Product code: **ODP** · Orthopedic  
Source: <https://www.510kdatabase.net/k082309/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Cervical (ODP)
Date received	Aug 13, 2008
Decision date	Apr 9, 2009
Days to decision	239 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Seaspine, Inc.</b>
Location	Vista, CA, US
Contact	JEFF BRITTAN
510(k) history	27 submissions · 27 cleared · 2005-2023

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k082309/> 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 19, 2026