

**K100698 STCC**Mar 14, 2011  
368 days to decisionK100698 · Product code: **ODP** · Orthopedic  
Source: <https://www.510kdatabase.net/k100698/>**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	Mar 11, 2010
Decision date	Mar 14, 2011
Days to decision	368 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Cardinal Spine, LLC</b>
Location	San Diego, CA, US
Contact	DAVID J COLLETTE
510(k) history	4 submissions · 4 cleared · 2011-2016

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k100698/> 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 17, 2026