

**K120275 SYNTHES ACIS/VERTEBRAL SPACER CR**May 25, 2012  
116 days to decisionK120275 · Product code: **ODP** · Orthopedic  
Source: <https://www.510kdatabase.net/k120275/>**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	Jan 30, 2012
Decision date	May 25, 2012
Days to decision	116 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Synthes (Usa), LLC</b>
Location	19380, PA, US
Contact	HEATHER GUERIN
510(k) history	7 submissions · 7 cleared · 2010-2018

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k120275/> 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 June 20, 2026