

**K030866 SYNTHES ANTERIOR CSLP SYSTEM**Apr 18, 2003  
30 days to decisionK030866 · Product code: **KWQ** · Orthopedic  
Source: <https://www.510kdatabase.net/k030866/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Intervertebral Body (KWQ)
Date received	Mar 19, 2003
Decision date	Apr 18, 2003
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Synthes (Usa)</b>
Location	Mchenry, IL, US
Contact	JONATHAN GILBERT
510(k) history	411 submissions · 394 cleared · 1977-2015

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k030866/> 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 14, 2026