

**K000742 SYNTHES ANTERIOR CSLP SYSTEM**Mar 29, 2000  
22 days to decisionK000742 · Product code: **KWQ** · Orthopedic  
Source: <https://www.510kdatabase.net/k000742/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Appliance, Fixation, Spinal Intervertebral Body (KWQ)
Date received	Mar 7, 2000
Decision date	Mar 29, 2000
Days to decision	22 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Synthes Spine</b>
Location	Paoli, PA, US
Contact	JONATHAN GILBERT
510(k) history	36 submissions · 32 cleared · 1995-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k000742/> 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 21, 2026