

**K925351 SYNTHES(U.S.A.) ANTERIOR SPINAL PLATE**Nov 5, 1993  
379 days to decisionK925351 · Product code: **KWQ** · Orthopedic  
Source: <https://www.510kdatabase.net/k925351/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Intervertebral Body (KWQ)
Date received	Oct 22, 1992
Decision date	Nov 5, 1993
Days to decision	379 days
Third-party review	No

**APPLICANT**

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Company	<b>Synthes (Usa)</b>
Location	Mchenry, IL, US
Contact	DIANE C TIERNAN
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/k925351/>, 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