

**K973868 SSCS (SEGMENTAL SPINAL CORRECTION SYSTEM)**Jan 8, 1998  
90 days to decisionK973868 · Product code: **KWP** · Orthopedic  
Source: <https://www.510kdatabase.net/k973868/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Interlaminar (KWP)
Date received	Oct 10, 1997
Decision date	Jan 8, 1998
Days to decision	90 days
Third-party review	No

**APPLICANT**

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Company	<b>Osteotech, Inc.</b>
Location	San Mateo, CA, US
Contact	CHRISTOPHER TALBOT
510(k) history	24 submissions · 21 cleared · 1985-2008

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k973868/>; 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