

**K010515 MODIFICATION TO SYNERGY SPINAL SYSTEM**Mar 22, 2001  
28 days to decisionK010515 · Product code: **KWP** · Orthopedic  
Source: <https://www.510kdatabase.net/k010515/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Appliance, Fixation, Spinal Interlaminar (KWP)
Date received	Feb 22, 2001
Decision date	Mar 22, 2001
Days to decision	28 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Interpore Cross Intl.</b>
Location	Irvine, CA, US
Contact	LYNN M RODARTI
510(k) history	39 submissions · 38 cleared · 1998-2005

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k010515/> 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 20, 2026