

**K051716 VERTEBRON PSS PEDICLE SCREW SYSTEM**Sep 7, 2005  
72 days to decisionK051716 · Product code: **KWP** · Orthopedic  
Source: <https://www.510kdatabase.net/k051716/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Appliance, Fixation, Spinal Interlaminar (KWP)
Date received	Jun 27, 2005
Decision date	Sep 7, 2005
Days to decision	72 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Vertebtron, Inc.</b>
Location	Stratford, CT, US
Contact	LUIS NESPRIDO
510(k) history	11 submissions · 11 cleared · 2004-2008

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k051716/> 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 19, 2026