

**K122509 VERTIFLEX SPINOUS PROCESS FIXATION PLATE**Dec 17, 2012  
122 days to decisionK122509 · Product code: **KWP** · Orthopedic  
Source: <https://www.510kdatabase.net/k122509/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Interlaminar (KWP)
Date received	Aug 17, 2012
Decision date	Dec 17, 2012
Days to decision	122 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Vertiflex, Inc.</b>
Location	San Clemente, CA, US
Contact	STEVE REITZLER
510(k) history	5 submissions · 5 cleared · 2008-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k122509/> 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 18, 2026