

**K014090 ENDIUS SPINAL FIXATION SYSTEM**Mar 11, 2002  
90 days to decisionK014090 · Product code: **KWP** · Orthopedic  
Source: <https://www.510kdatabase.net/k014090/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Interlaminar (KWP)
Date received	Dec 11, 2001
Decision date	Mar 11, 2002
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Endius, Inc.</b>
Location	Plainville, MA, US
Contact	GENE DIPOTO
510(k) history	33 submissions · 33 cleared · 1997-2008

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k014090/> 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