

**K092656 LANX POSTERIOR CERVICOTHORACIC SPINAL  
FIXATION SYSTEM**Nov 24, 2009  
88 days to decisionK092656 · Product code: **KWP** · Orthopedic  
Source: <https://www.510kdatabase.net/k092656/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Interlaminar (KWP)
Date received	Aug 28, 2009
Decision date	Nov 24, 2009
Days to decision	88 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Lanx, Inc.</b>
Location	Broomfield, CO, US
Contact	ANDREW LAMBORNE
510(k) history	23 submissions · 23 cleared · 2009-2013

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k092656/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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