

**K071905 POSTERIOR CERVICOTHORACIC SPINAL FIXATION SYSTEM (PCFS)**Dec 28, 2007  
171 days to decisionK071905 · Product code: **KWQ** · Orthopedic  
Source: <https://www.510kdatabase.net/k071905/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Intervertebral Body (KWQ)
Date received	Jul 10, 2007
Decision date	Dec 28, 2007
Days to decision	171 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Lanx, LLC</b>
Location	Pleasant Hill, CA, US
Contact	JANICE M HOGAN
510(k) history	13 submissions · 13 cleared · 2004-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k071905/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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