

**K103660 LANX CERVICAL INTERVERTEBRAL BODY FUSION SYSTEM**Feb 28, 2011  
75 days to decisionK103660 · Product code: **ODP** · Orthopedic  
Source: <https://www.510kdatabase.net/k103660/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Intervertebral Fusion Device With Bone Graft, Cervical (ODP)
Date received	Dec 15, 2010
Decision date	Feb 28, 2011
Days to decision	75 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Lanx, Inc.</b>
Location	Broomfield, CO, US
Contact	WILLIAM SANDUL
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/k103660/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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