

**K103237 LANX SPINAL SYSTEM**Mar 25, 2011  
143 days to decisionK103237 · Product code: **NKB** · Orthopedic  
Source: <https://www.510kdatabase.net/k103237/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Thoracolumbosacral Pedicle Screw System (NKB)
Date received	Nov 2, 2010
Decision date	Mar 25, 2011
Days to decision	143 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k103237/> 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