

**K103040 LANX POSTERIOR CERVICOTHORACIC SPINAL
FIXATION SYSTEM (PCFS)**Feb 8, 2011
118 days to decisionK103040 · Product code: **NKB** · Orthopedic
Source: <https://www.510kdatabase.net/k103040/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Thoracolumbosacral Pedicle Screw System (NKB)
Date received	Oct 13, 2010
Decision date	Feb 8, 2011
Days to decision	118 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Lanx, LLC
Location	Pleasant Hill, CA, US
Contact	DAMON BELLONI
510(k) history	13 submissions · 13 cleared · 2004-2012

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k103040/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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