FDA 510(k) Premarket Notification

K103237 LANX SPINAL SYSTEM

CLEARED

Mar 25, 2011
143 days to decision

K103237 · 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	**Lanx, Inc.**
Location	Broomfield, CO, US
Contact	DAMON BELLONI
510(k) history	23 submissions · 23 cleared · 2009-2013