FDA 510(k) Premarket Notification

K092656 LANX POSTERIOR CERVICOTHORACIC SPINAL FIXATION SYSTEM

CLEARED

Nov 24, 2009
88 days to decision

K092656 · Product code: **KWP** · Orthopedic
Source: https://www.510kdatabase.net/k092656/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Interlaminal (KWP)
Date received	Aug 28, 2009
Decision date	Nov 24, 2009
Days to decision	88 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Lanx, Inc.**
Location	Broomfield, CO, US
Contact	ANDREW LAMBORNE
510(k) history	23 submissions · 23 cleared · 2009-2013