FDA 510(k) Premarket Notification

CLEARED

K090252 MODIFICATION TO: LANX SPINAL FIXATION SYSTEM

Mar 4, 2009
30 days to decision

K090252 · Product code: **KWP** · Orthopedic
Source: https://www.510kdatabase.net/k090252/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Appliance, Fixation, Spinal Interlaminal (KWP)
Date received	Feb 2, 2009
Decision date	Mar 4, 2009
Days to decision	30 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