

K090252 MODIFICATION TO: LANX SPINAL FIXATION SYSTEMMar 4, 2009
30 days to decisionK090252 · Product code: **KWP** · Orthopedic
Source: <https://www.510kdatabase.net/k090252/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Appliance, Fixation, Spinal Interlaminar (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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k090252/> Data sourced from FDA 510(k) public records (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. - Generated May 18, 2026