

K090316 MODIFICATION TO: LANX ANTERIOR CERVICAL PLATE SYSTEMApr 7, 2009
57 days to decisionK090316 · Product code: **KWQ** · Orthopedic
Source: <https://www.510kdatabase.net/k090316/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Appliance, Fixation, Spinal Intervertebral Body (KWQ)
Date received	Feb 9, 2009
Decision date	Apr 7, 2009
Days to decision	57 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/k090316/> 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