FDA 510(k) Premarket Notification

K090316 MODIFICATION TO: LANX ANTERIOR CERVICAL PLATE SYSTEM

CLEARED

Apr 7, 2009
57 days to decision

K090316 · 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