

**K082774 LANX CERVICAL INTERVERTEBRAL BODY FUSION
DEVICE**

Oct 21, 2008
29 days to decision

K082774 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k082774/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Sep 22, 2008
Decision date	Oct 21, 2008
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Lanx, LLC
Location	Pleasant Hill, CA, US
Contact	Alan Burkholder
510(k) history	13 submissions · 13 cleared · 2004-2012

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k082774/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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