FDA 510(k) Premarket Notification

CLEARED

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