FDA 510(k) Premarket Notification

K123752 ACCULIF TL AND PL CAGE

CLEARED

Jan 14, 2013
39 days to decision

K123752 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k123752/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Dec 6, 2012
Decision date	Jan 14, 2013
Days to decision	39 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Coalign Innovations, Inc.**
Location	Washington, DC, US
Contact	JUSTIN EGGLETON
510(k) history	9 submissions · 9 cleared · 2011-2013