FDA 510(k) Premarket Notification

K113465 ACCULIF TL CAGE

CLEARED

Dec 12, 2011
20 days to decision

K113465 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k113465/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Nov 22, 2011
Decision date	Dec 12, 2011
Days to decision	20 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