FDA 510(k) Premarket Notification

K141942 MIDLINE II TI

CLEARED

Nov 7, 2014
113 days to decision

K141942 · Product code: **OVD** · Orthopedic
Source: https://www.510kdatabase.net/k141942/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Integrated Fixation, Lumbar (OVD)
Date received	Jul 17, 2014
Decision date	Nov 7, 2014
Days to decision	113 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Centinel Spine, Inc.**
Location	West Chester, PA, US
Contact	JUSTIN EGGLETON
510(k) history	10 submissions · 10 cleared · 2012-2018