FDA 510(k) Premarket Notification

K100698 STCC

CLEARED

Mar 14, 2011
368 days to decision

K100698 · Product code: **ODP** · Orthopedic
Source: https://www.510kdatabase.net/k100698/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Cervical (ODP)
Date received	Mar 11, 2010
Decision date	Mar 14, 2011
Days to decision	368 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Cardinal Spine, LLC**
Location	San Diego, CA, US
Contact	DAVID J COLLETTE
510(k) history	4 submissions · 4 cleared · 2011-2016