FDA 510(k) Premarket Notification

K082309 CAMBRIA

CLEARED

Apr 9, 2009
239 days to decision

K082309 · Product code: **ODP** · Orthopedic
Source: https://www.510kdatabase.net/k082309/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Cervical (ODP)
Date received	Aug 13, 2008
Decision date	Apr 9, 2009
Days to decision	239 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Seaspine, Inc.**
Location	Vista, CA, US
Contact	JEFF BRITTAN
510(k) history	27 submissions · 27 cleared · 2005-2023