FDA 510(k) Premarket Notification

K120293 SPINEOLOGY PEEK LUMBAR INTERBODY FUSION DRIVE

CLEARED

Feb 29, 2012
29 days to decision

K120293 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k120293/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Jan 31, 2012
Decision date	Feb 29, 2012
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Spineology, Inc.**
Location	Stillwater, MN, US
Contact	BRYAN BECKER
510(k) history	54 submissions · 51 cleared · 1999-2025