FDA 510(k) Premarket Notification

K110933 SPINEOLOGY PEEK LUMBAR INTERBODY FUSION DEVICE

CLEARED

Jun 13, 2011
70 days to decision

K110933 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k110933/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Apr 4, 2011
Decision date	Jun 13, 2011
Days to decision	70 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