

**K123652 SPINEOLOGY PEEK LUMBAR INTERBODY FUSION
DEVICE**Mar 22, 2013
115 days to decisionK123652 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k123652/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Nov 27, 2012
Decision date	Mar 22, 2013
Days to decision	115 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k123652/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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