

K182322 Duo Lumbar Interbody Fusion DeviceOct 24, 2018
58 days to decisionK182322 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k182322/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Aug 27, 2018
Decision date	Oct 24, 2018
Days to decision	58 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

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