

K200958 MODULIF-A Anterior Lumbar Interbody Fusion SystemAug 28, 2020
140 days to decisionK200958 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k200958/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Apr 10, 2020
Decision date	Aug 28, 2020
Days to decision	140 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Spinal Stability, LLC
Location	Austin, TX, US
Contact	David Krueger
510(k) history	1 submissions · 1 cleared · 2020-2020

REGULATORY CONSULTANT

Consulting firm	The OrthoMedix Group, Inc.
Contact	J.D. Webb

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov

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