

K101301 STALIF MIDLINESep 24, 2010
137 days to decisionK101301 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k101301/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	May 10, 2010
Decision date	Sep 24, 2010
Days to decision	137 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Centinel Spine
Location	Washington, DC, US
Contact	JUSTIN EGGLETON
510(k) history	1 submissions · 1 cleared · 2010-2010

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