

**K160663 Lumbar Interbody Fusion System (ALIF, PLIF, LLIF, TPLIF)**Nov 16, 2016  
252 days to decisionK160663 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k160663/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Mar 9, 2016
Decision date	Nov 16, 2016
Days to decision	252 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Evolution Spine, LLC</b>
Location	Dallas, TX, US
Contact	Ashton Kouzbari
510(k) history	5 submissions · 5 cleared · 2016-2019

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k160663/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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