

**K160874 Nanovis Intervertebral Body Fusion System and FortiCore®**Jun 28, 2016  
90 days to decisionK160874 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k160874/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Mar 30, 2016
Decision date	Jun 28, 2016
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Nanovis, LLC</b>
Location	San Diego, CA, US
Contact	Matthew Hedrick
510(k) history	10 submissions · 10 cleared · 2011-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k160874/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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