

**K162715 SeaSpine Spacer System- Hollywood NanoMetalene,  
Hollywood VI NanoMetalene, Ventura NanoMetalene**Oct 28, 2016  
29 days to decisionK162715 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k162715/>**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	Sep 29, 2016
Decision date	Oct 28, 2016
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary
Other names	SeaSpine Cambria NanoMetalene

**APPLICANT**

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Company	<b>SeaSpine Orthopedics Corporation</b>
Location	Carlsbad, CA, US
Contact	Gina Flores
510(k) history	66 submissions · 66 cleared · 2016-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k162715/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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