

**K141896 COROENT SYSTEM**Nov 20, 2014  
129 days to decisionK141896 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k141896/>**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	Jul 14, 2014
Decision date	Nov 20, 2014
Days to decision	129 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Nu Vasive, Incorporated</b>
Location	San Diego, CA, US
Contact	OLGA LEWIS
510(k) history	112 submissions · 112 cleared · 2012-2023

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k141896/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 17, 2026