

**K143595 NuVasive(R) CCX-L Interbody System**Apr 29, 2015  
132 days to decisionK143595 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k143595/>**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	Dec 18, 2014
Decision date	Apr 29, 2015
Days to decision	132 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	Cynthia Adams
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/k143595/> 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