

**K200953 NuVasive® Cohere® Thoracolumbar Interbody System**Oct 6, 2020  
180 days to decisionK200953 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k200953/>**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	Apr 9, 2020
Decision date	Oct 6, 2020
Days to decision	180 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	Michelle Cheung
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/k200953/> 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 15, 2026