

K210439 NuVasive MOD-EX XLIF Interbody SystemMay 12, 2021
89 days to decisionK210439 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k210439/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Feb 12, 2021
Decision date	May 12, 2021
Days to decision	89 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Nu Vasive, Incorporated
Location	San Diego, CA, US
Contact	Manthan J. Damani
510(k) history	112 submissions · 112 cleared · 2012-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k210439/> Data sourced from FDA 510(k) public records (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. - Generated May 13, 2026