

K153105 MLX™ - Medial Lateral Expandable Lumbar Interbody SystemJul 11, 2016
258 days to decisionK153105 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k153105/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Oct 27, 2015
Decision date	Jul 11, 2016
Days to decision	258 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Nu Vasive, Incorporated
Location	San Diego, CA, US
Contact	Olga Lewis
510(k) history	112 submissions · 112 cleared · 2012-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k153105/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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