

**K140770 MLX - MEDIAL LATERAL EXPANDABLE LUMBAR INTERBODY SYSTEM**Jul 25, 2014  
120 days to decisionK140770 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k140770/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Mar 27, 2014
Decision date	Jul 25, 2014
Days to decision	120 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k140770/>. 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 17, 2026