FDA 510(k) Premarket Notification

# K151374 NuVasive MLX-Medial Lateral Expandable Lumbar Interbody System, NuVasive AP Expandable XLIF System

**CLEARED**

Aug 6, 2015
76 days to decision

K151374 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k151374/

## 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 | May 22, 2015 |
| Decision date | Aug 6, 2015 |
| Days to decision | 76 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.net
Device record: https://www.510kdatabase.net/k151374/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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