FDA 510(k) Premarket Notification

# K172123 NuVasive® Modulus XLIF Interbody System

**CLEARED**

Oct 11, 2017
89 days to decision

K172123 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k172123/

## 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 | Jul 14, 2017 |
| Decision date | Oct 11, 2017 |
| Days to decision | 89 days |
| Third-party review | No |
| Summary / Statement | Summary |

## APPLICANT

| | |
|---|---|
| Company | **Nu Vasive, Incorporated** |
| Location | San Diego, CA, US |
| Contact | Cynthia Adams |
| 510(k) history | 112 submissions · 112 cleared · 2012-2023 |