

K172123 NuVasive® Modulus XLIF Interbody SystemOct 11, 2017
89 days to decisionK172123 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k172123/>**SUBMISSION DETAILS**

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

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