

K172341 NuVasive® Modulus TLIF Interbody SystemOct 26, 2017
85 days to decisionK172341 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k172341/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Aug 2, 2017
Decision date	Oct 26, 2017
Days to decision	85 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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