

K230894 NuVasive Modulus ALIF SystemJun 16, 2023
77 days to decisionK230894 · Product code: **OVD** · Orthopedic
Source: <https://www.510kdatabase.net/k230894/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Integrated Fixation, Lumbar (OVD)
Date received	Mar 31, 2023
Decision date	Jun 16, 2023
Days to decision	77 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Nu Vasive, Incorporated
Location	San Diego, CA, US
Contact	Alexander Stevens
510(k) history	112 submissions · 112 cleared · 2012-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k230894/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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