FDA 510(k) Premarket Notification

K230894 NuVasive Modulus ALIF System

CLEARED

Jun 16, 2023
77 days to decision

K230894 · 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