FDA 510(k) Premarket Notification

K201692 NuVasive Modulus XLIF Interbody System

CLEARED

Oct 29, 2020
129 days to decision

K201692 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k201692/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Jun 22, 2020
Decision date	Oct 29, 2020
Days to decision	129 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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