FDA 510(k) Premarket Notification

CLEARED

K172341 NuVasive® Modulus TLIF Interbody System

Oct 26, 2017
85 days to decision

K172341 · 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