FDA 510(k) Premarket Notification

K171633 NuVasive TLX Interbody System

CLEARED

Sep 28, 2017
118 days to decision

K171633 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k171633/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Jun 2, 2017
Decision date	Sep 28, 2017
Days to decision	118 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