FDA 510(k) Premarket Notification

CLEARED

K173892 NuVasive® XLX Interbody System

May 4, 2018
134 days to decision

K173892 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k173892/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Dec 21, 2017
Decision date	May 4, 2018
Days to decision	134 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