FDA 510(k) Premarket Notification

K182974 NuVasive Reline System

CLEARED

Feb 13, 2019
110 days to decision

K182974 · Product code: **NKB** · Orthopedic
Source: https://www.510kdatabase.net/k182974/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Thoracolumbosacral Pedicle Screw System (NKB)
Date received	Oct 26, 2018
Decision date	Feb 13, 2019
Days to decision	110 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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