FDA 510(k) Premarket Notification

CLEARED

K180198 NuVasive® VuePoint® II OCT System

Mar 15, 2018
50 days to decision

K180198 · Product code: **NKG** · Orthopedic
Source: https://www.510kdatabase.net/k180198/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Posterior Cervical Screw System (NKG)
Date received	Jan 24, 2018
Decision date	Mar 15, 2018
Days to decision	50 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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