

K180198 NuVasive® VuePoint® II OCT SystemMar 15, 2018
50 days to decisionK180198 · 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

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