FDA 510(k) Premarket Notification

K213654 NuVasive Reline Cervical System

CLEARED

Feb 23, 2022
96 days to decision

K213654 · Product code: **NKG** · Orthopedic
Source: https://www.510kdatabase.net/k213654/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Posterior Cervical Screw System (NKG)
Date received	Nov 19, 2021
Decision date	Feb 23, 2022
Days to decision	96 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

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