FDA 510(k) Premarket Notification

K180550 NuVasive Monolith Cervical Corpectomy System

CLEARED

Nov 20, 2018
264 days to decision

K180550 · Product code: **PLR** · Orthopedic
Source: https://www.510kdatabase.net/k180550/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spinal Vertebral Body Replacement Device - Cervical (PLR)
Date received	Mar 1, 2018
Decision date	Nov 20, 2018
Days to decision	264 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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