FDA 510(k) Premarket Notification

K200953 NuVasive® Cohere® Thoracolumbar Interbody System

CLEARED

Oct 6, 2020
180 days to decision

K200953 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k200953/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Apr 9, 2020
Decision date	Oct 6, 2020
Days to decision	180 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