FDA 510(k) Premarket Notification

CLEARED

K173030 Vertera Spine Cohere Cervical Interbody Fusion Device

Nov 21, 2017
54 days to decision

K173030 · Product code: **ODP** · Orthopedic
Source: https://www.510kdatabase.net/k173030/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Cervical (ODP)
Date received	Sep 28, 2017
Decision date	Nov 21, 2017
Days to decision	54 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