

K173030 Vertera Spine Cohere Cervical Interbody Fusion DeviceNov 21, 2017
54 days to decisionK173030 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k173030/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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