

K172676 NuVasive® Modulus-C Interbody SystemDec 14, 2017
100 days to decisionK172676 · Product code: **ODP** · Orthopedic
Source: <https://www.510kdatabase.net/k172676/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Cervical (ODP)
Date received	Sep 5, 2017
Decision date	Dec 14, 2017
Days to decision	100 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Nu Vasive, Incorporated
Location	San Diego, CA, US
Contact	Marysa E. Loustalot
510(k) history	112 submissions · 112 cleared · 2012-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k172676/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 16, 2026