

K180550 NuVasive Monolith Cervical Corpectomy SystemNov 20, 2018
264 days to decisionK180550 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k180550/> 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