

**K173606 SeaSpine Vu a•POD Prime NanoMetalene
Intervertebral Body Fusion Device, SeaSpine Vu a•POD Prime
Intervertebral Body Fusion Device**Apr 13, 2018
143 days to decisionK173606 · Product code: **OVD** · Orthopedic
Source: <https://www.510kdatabase.net/k173606/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Integrated Fixation, Lumbar (OVD)
Date received	Nov 21, 2017
Decision date	Apr 13, 2018
Days to decision	143 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	SeaSpine Orthopedics Corporation
Location	Carlsbad, CA, US
Contact	Gina Flores
510(k) history	66 submissions · 66 cleared · 2016-2025

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k173606/> 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 15, 2026