

K172253 Altus Spine Cervical Interbody Fusion SystemNov 21, 2017
118 days to decisionK172253 · Product code: **ODP** · Orthopedic
Source: <https://www.510kdatabase.net/k172253/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Cervical (ODP)
Date received	Jul 26, 2017
Decision date	Nov 21, 2017
Days to decision	118 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Altus Partners, LLC
Location	Newtown Square, PA, US
Contact	Mark Melton
510(k) history	17 submissions · 17 cleared · 2015-2025

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k172253/> 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 26, 2026