

K200922 Altus Spine HA Pedicle Screw SystemJan 26, 2021
295 days to decisionK200922 · Product code: **NKB** · Orthopedic
Source: <https://www.510kdatabase.net/k200922/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Thoracolumbosacral Pedicle Screw System (NKB)
Date received	Apr 6, 2020
Decision date	Jan 26, 2021
Days to decision	295 days
Third-party review	No
Combination product	No
PCCP authorized	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/k200922/> 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