

**K170942 The Thunderbolt™ Minimally Invasive and Lancer™
Open Pedicle Screw Systems**

Apr 26, 2017
27 days to decision

K170942 · Product code: **NKB** · Orthopedic
Source: <https://www.510kdatabase.net/k170942/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Thoracolumbosacral Pedicle Screw System (NKB)
Date received	Mar 30, 2017
Decision date	Apr 26, 2017
Days to decision	27 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Choicespine, LP
Location	Chesterland, OH, US
Contact	Kim Finch
510(k) history	22 submissions · 22 cleared · 2008-2019

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k170942/> 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 June 20, 2026