

**K171456 Thunderbolt™ Minimally Invasive and Lancer™ Open Pedicle Screw Systems**

Aug 3, 2017  
78 days to decision

K171456 · Product code: **NKB** · Orthopedic  
Source: <https://www.510kdatabase.net/k171456/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Thoracolumbosacral Pedicle Screw System (NKB)
Date received	May 17, 2017
Decision date	Aug 3, 2017
Days to decision	78 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Choicespine</b>
Location	Knoxville, TN, US
Contact	Kim Finch
510(k) history	4 submissions · 4 cleared · 2014-2019

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k171456/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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