

K170821 ChoiceSpine™ Proliant® Posterior Pedicle Screw and Hook Fixation SystemJun 8, 2017
80 days to decisionK170821 · Product code: **NKB** · Orthopedic
Source: <https://www.510kdatabase.net/k170821/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Thoracolumbosacral Pedicle Screw System (NKB)
Date received	Mar 20, 2017
Decision date	Jun 8, 2017
Days to decision	80 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.netDevice record: <https://www.510kdatabase.net/k170821/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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