

K132049 THUNDERBOLT AND LANCER PEDICLE SCREW SYSTEMSAug 15, 2014
409 days to decisionK132049 · Product code: **NKB** · Orthopedic
Source: <https://www.510kdatabase.net/k132049/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Thoracolumbosacral Pedicle Screw System (NKB)
Date received	Jul 2, 2013
Decision date	Aug 15, 2014
Days to decision	409 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Choicespine
Location	Knoxville, TN, US
Contact	KIM FINCH
510(k) history	4 submissions · 4 cleared · 2014-2019

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k132049/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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