

**K082032 SEQUOIA SPINAL SYSTEM (MODEL 3306),
SPEEDLINK TRANSVERSE CONNECTOR (MODELS 3308, 3309,
3310)**Oct 6, 2008
81 days to decisionK082032 · Product code: **NKB** · Orthopedic
Source: <https://www.510kdatabase.net/k082032/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Thoracolumbosacral Pedicle Screw System (NKB)
Date received	Jul 17, 2008
Decision date	Oct 6, 2008
Days to decision	81 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Abbott Spine, Inc.
Location	Austin, TX, US
Contact	DAVID PADGETT
510(k) history	13 submissions · 13 cleared · 2005-2009

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k082032/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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