

**K103383 SPARTEK VARIABLE ANGLE PEDICLE SCREW
POSTERIOR FUSION SYSTEM**Feb 17, 2011
91 days to decisionK103383 · Product code: **NKB** · Orthopedic
Source: <https://www.510kdatabase.net/k103383/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Thoracolumbosacral Pedicle Screw System (NKB)
Date received	Nov 18, 2010
Decision date	Feb 17, 2011
Days to decision	91 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Spartek Medical, Inc.
Location	Alameda, CA, US
Contact	HENRY KLYCE
510(k) history	2 submissions · 2 cleared · 2010-2011

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k103383/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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