

**K072183 ST360 SPINAL FIXATION SYSTEM, MODEL
07.0029-7500.5600**

Sep 5, 2007
30 days to decision

K072183 · Product code: **NKB** · Orthopedic
Source: <https://www.510kdatabase.net/k072183/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Thoracolumbosacral Pedicle Screw System (NKB)
Date received	Aug 6, 2007
Decision date	Sep 5, 2007
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Zimmer Spine, Inc.
Location	Minneapolis, MN, US
Contact	TIM CRABTREE
510(k) history	38 submissions · 35 cleared · 2004-2016

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k072183/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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