FDA 510(k) Premarket Notification

K132884 PATHFINDER NXT MINIMALLY INVASIVE PEDICLE SCREW SYSTEM

CLEARED

Nov 25, 2013
73 days to decision

K132884 · Product code: **NKB** · Orthopedic
Source: https://www.510kdatabase.net/k132884/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Thoracolumbosacral Pedicle Screw System (NKB)
Date received	Sep 13, 2013
Decision date	Nov 25, 2013
Days to decision	73 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k132884/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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