FDA 510(k) Premarket Notification

K121671 PATHFINDER NXT MINIMALLY INVASIVE PEDICLE SCREW SYSTEM

CLEARED

Jul 18, 2012
42 days to decision

K121671 · Product code: **NKB** · Orthopedic
Source: https://www.510kdatabase.net/k121671/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Thoracolumbosacral Pedicle Screw System (NKB)
Date received	Jun 6, 2012
Decision date	Jul 18, 2012
Days to decision	42 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k121671/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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