FDA 510(k) Premarket Notification

K042702 SILHOUETTE SPINAL FIXATION SYSTEM

CLEARED

Nov 23, 2004
54 days to decision

K042702 · Product code: **NKB** · Orthopedic
Source: https://www.510kdatabase.net/k042702/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Thoracolumbosacral Pedicle Screw System (NKB)
Date received	Sep 30, 2004
Decision date	Nov 23, 2004
Days to decision	54 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/k042702/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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