FDA 510(k) Premarket Notification

K072672 SEQUOIA SPINAL SYSTEM, MODEL 3300 SERIES

CLEARED

Nov 28, 2007
68 days to decision

K072672 · Product code: **NKB** · Orthopedic
Source: https://www.510kdatabase.net/k072672/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Thoracolumbosacral Pedicle Screw System (NKB)
Date received	Sep 21, 2007
Decision date	Nov 28, 2007
Days to decision	68 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Abbott Spine, Inc.**
Location	Austin, TX, US
Contact	DAVID PADGETT
510(k) history	13 submissions · 13 cleared · 2005-2009