FDA 510(k) Premarket Notification

CLEARED

K102204 CHOICE SPINE STARFIRE PEDICLE SCREW SYSTEM

Nov 15, 2010
102 days to decision

K102204 · Product code: **NKB** · Orthopedic
Source: https://www.510kdatabase.net/k102204/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Thoracolumbosacral Pedicle Screw System (NKB)
Date received	Aug 5, 2010
Decision date	Nov 15, 2010
Days to decision	102 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Choicespine, LP**
Location	Chesterland, OH, US
Contact	TODD HAWKINS
510(k) history	22 submissions · 22 cleared · 2008-2019

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k102204/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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