FDA 510(k) Premarket Notification

CLEARED

K171456 Thunderbolt™ Minimally Invasive and Lancer™ Open Pedicle Screw Systems

Aug 3, 2017
78 days to decision

K171456 · Product code: **NKB** · Orthopedic
Source: https://www.510kdatabase.net/k171456/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Thoracolumbosacral Pedicle Screw System (NKB)
Date received	May 17, 2017
Decision date	Aug 3, 2017
Days to decision	78 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Choicespine**
Location	Knoxville, TN, US
Contact	Kim Finch
510(k) history	4 submissions · 4 cleared · 2014-2019