FDA 510(k) Premarket Notification

K082592 LDR SPINE EASYSPINE SYSTEM

CLEARED

Oct 8, 2008
30 days to decision

K082592 · Product code: **NKB** · Orthopedic
Source: https://www.510kdatabase.net/k082592/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Thoracolumbosacral Pedicle Screw System (NKB)
Date received	Sep 8, 2008
Decision date	Oct 8, 2008
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Ldr Spine USA**
Location	Austin, TX, US
Contact	NOAH BARTSCH
510(k) history	25 submissions · 25 cleared · 2005-2016