

**K121103 LDR SPINE USA SPINE TUNE, TL SPINAL SYSTEM,  
LDR SPINE USA EASYSPINE, POSTERIOR SPINAL SYSTEM,  
LDR SPINE USA MC IMPLANT S**

Aug 24, 2012  
135 days to decision

K121103 · Product code: **NKB** · Orthopedic  
Source: <https://www.510kdatabase.net/k121103/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Thoracolumbosacral Pedicle Screw System (NKB)
Date received	Apr 11, 2012
Decision date	Aug 24, 2012
Days to decision	135 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Ldr Spine USA, Inc.</b>
Location	Austin, TX, US
Contact	BRADLEY W STRASSER
510(k) history	10 submissions · 10 cleared · 2011-2017

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k121103/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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