

**K082592 LDR SPINE EASYSPINE SYSTEM**Oct 8, 2008  
30 days to decisionK082592 · Product code: **NKB** · Orthopedic  
Source: <https://www.510kdatabase.net/k082592/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k082592/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 17, 2026