

**K141873 RANGE/MESA/DENALI SPINAL SYSTEM**

Sep 8, 2014  
59 days to decision

K141873 · Product code: **NKB** · Orthopedic  
Source: <https://www.510kdatabase.net/k141873/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Thoracolumbosacral Pedicle Screw System (NKB)
Date received	Jul 11, 2014
Decision date	Sep 8, 2014
Days to decision	59 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>K2m, Inc.</b>
Location	Leesburg, VA, US
Contact	NANCY GLEZEN
510(k) history	100 submissions · 97 cleared · 2007-2026

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

Device record: <https://www.510kdatabase.net/k141873/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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