

**K121551 RESOLUTE FACET SCREW SYSTEM**Oct 23, 2012  
151 days to decisionK121551 · Product code: **MRW** · Orthopedic  
Source: <https://www.510kdatabase.net/k121551/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Facet Screw Spinal Device (MRW)
Date received	May 25, 2012
Decision date	Oct 23, 2012
Days to decision	151 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Neurostructures, LLC</b>
Location	Colorado Springs, CO, US
Contact	MEREDITH MAY
510(k) history	3 submissions · 3 cleared · 2012-2013

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k121551/> 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 18, 2026