

**K120340 VENUS FACET SCREW SYSTEM**Oct 19, 2012  
259 days to decisionK120340 · Product code: **MRW** · Orthopedic  
Source: <https://www.510kdatabase.net/k120340/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Facet Screw Spinal Device (MRW)
Date received	Feb 3, 2012
Decision date	Oct 19, 2012
Days to decision	259 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Apollo Spine, Inc.</b>
Location	Newport Beach, CA, US
Contact	CHRISTINE SANTAGATE
510(k) history	6 submissions · 6 cleared · 2009-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k120340/> 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