

**K070922 MODIFICATION TO SPINAL USA VBR SYSTEM**Sep 19, 2007  
170 days to decisionK070922 · Product code: **KWQ** · Orthopedic  
Source: <https://www.510kdatabase.net/k070922/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Intervertebral Body (KWQ)
Date received	Apr 2, 2007
Decision date	Sep 19, 2007
Days to decision	170 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Spinal USA</b>
Location	Brandon, MS, US
Contact	JEFFREY JOHNSON
510(k) history	23 submissions · 23 cleared · 2006-2013

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k070922/> 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