

**K133575 SPINEVISION LUMIS CANNULATED PEDICLE SCREW  
FIXATION SYSTEM, SPINEVISION U.L.I.S. PEDICLE SCREW  
FIXATION SYSTEM**Jun 30, 2014  
222 days to decisionK133575 · Product code: **MNH** · Orthopedic  
Source: <https://www.510kdatabase.net/k133575/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Orthosis, Spondylolisthesis Spinal Fixation (MNH)
Date received	Nov 20, 2013
Decision date	Jun 30, 2014
Days to decision	222 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Spinevision S.A.</b>
Location	Douglassville, PA, US
Contact	J.D. WEBB
510(k) history	5 submissions · 5 cleared · 2012-2016

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k133575/> 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