

**K132575 PASS LP SPINAL SYSTEM**Oct 3, 2013  
48 days to decisionK132575 · Product code: **OSH** · Orthopedic  
Source: <https://www.510kdatabase.net/k132575/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Pedicle Screw Spinal System, Adolescent Idiopathic Scoliosis (OSH)
Date received	Aug 16, 2013
Decision date	Oct 3, 2013
Days to decision	48 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Medicrea International</b>
Location	Round Rock, TX, US
Contact	AUDREY VION
510(k) history	17 submissions · 16 cleared · 2011-2022

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k132575/> 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 27, 2026