

**K963639 ACROMED PEDICLE SCREW**Nov 14, 1996  
63 days to decisionK963639 · Product code: **KWP** · Orthopedic  
Source: <https://www.510kdatabase.net/k963639/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Interlaminar (KWP)
Date received	Sep 12, 1996
Decision date	Nov 14, 1996
Days to decision	63 days
Third-party review	No

**APPLICANT**

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Company	<b>Acromed Corp.</b>
Location	Cleveland, OH, US
Contact	WILLIAM CHRISTIANSON
510(k) history	41 submissions · 22 cleared · 1984-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k963639/>; 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 21, 2026