

**K951116 ACROMED PEDICLE SCREW**Aug 11, 1995  
154 days to decisionK951116 · Product code: **MNH** · Orthopedic  
Source: <https://www.510kdatabase.net/k951116/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Orthosis, Spondylolisthesis Spinal Fixation (MNH)
Date received	Mar 10, 1995
Decision date	Aug 11, 1995
Days to decision	154 days
Third-party review	No

**APPLICANT**

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Company	<b>Acromed Corp.</b>
Location	Cleveland, OH, US
Contact	GREGORY CANNEDY
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/k951116/>; 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