

**K950999 KSF POSTERIOR SPINGAL FIXATOR FOR SEVERE SPONDYLOLISTHESIS**Jan 25, 1996  
328 days to decisionK950999 · Product code: **MNH** · Orthopedic  
Source: <https://www.510kdatabase.net/k950999/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Orthosis, Spondylolisthesis Spinal Fixation (MNH)
Date received	Mar 3, 1995
Decision date	Jan 25, 1996
Days to decision	328 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Buckman Co., Inc.</b>
Location	Mchenry, IL, US
Contact	DAVID W SCHLERF
510(k) history	111 submissions · 104 cleared · 1983-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k950999/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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