

**K020196 MODIFICATION TO SILHOUETTE SPINAL FIXATION SYSTEM**Feb 15, 2002  
24 days to decisionK020196 · Product code: **MNH** · Orthopedic  
Source: <https://www.510kdatabase.net/k020196/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Orthosis, Spondylolisthesis Spinal Fixation (MNH)
Date received	Jan 22, 2002
Decision date	Feb 15, 2002
Days to decision	24 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Sulzer Spine-Tech</b>
Location	Minneapolis, MN, US
Contact	KRISTYN M BENSON
510(k) history	7 submissions · 7 cleared · 1999-2002

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k020196/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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