

**K951846 MIRAGE SPINAL SYSTEM FOR SEVERE SPONDYLOLISTHESIS**Feb 14, 1996  
300 days to decisionK951846 · Product code: **KWP** · Orthopedic  
Source: <https://www.510kdatabase.net/k951846/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Interlaminar (KWP)
Date received	Apr 20, 1995
Decision date	Feb 14, 1996
Days to decision	300 days
Third-party review	No

**APPLICANT**

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Company	<b>Alphatec Mfg., Inc.</b>
Location	Palm Desert, CA, US
Contact	DAVID W SCHLERF
510(k) history	20 submissions · 18 cleared · 1990-2004

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k951846/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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