

**K022374 CADENCE SPINAL FIXATION SYSTEM**Sep 24, 2002  
64 days to decisionK022374 · Product code: **KWP** · Orthopedic  
Source: <https://www.510kdatabase.net/k022374/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Appliance, Fixation, Spinal Interlaminar (KWP)
Date received	Jul 22, 2002
Decision date	Sep 24, 2002
Days to decision	64 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/k022374/> 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 20, 2026