

**K983706 MODIFICATION OF TOWNLEY PEDICLE SCREW  
PLATING SYSTEM**Nov 12, 1998  
22 days to decisionK983706 · Product code: **KWP** · Orthopedic  
Source: <https://www.510kdatabase.net/k983706/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Appliance, Fixation, Spinal Interlaminar (KWP)
Date received	Oct 21, 1998
Decision date	Nov 12, 1998
Days to decision	22 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Sofamor Danek USA, Inc.</b>
Location	Memphis, TN, US
Contact	RICHARD W TREHARNE
510(k) history	41 submissions · 26 cleared · 1995-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k983706/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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