

**K123906 VERTEX RECONSTRUCTION SYSTEM, VERTEXT
SELECT RECONSTRUCTION SYSTEM**Apr 1, 2013
103 days to decisionK123906 · Product code: **KWP** · Orthopedic
Source: <https://www.510kdatabase.net/k123906/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Interlaminar (KWP)
Date received	Dec 19, 2012
Decision date	Apr 1, 2013
Days to decision	103 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Medtronic Sofamor Danek USA, Inc.
Location	Memphis, TN, US
Contact	CLAIRE EVANS
510(k) history	170 submissions · 159 cleared · 2000-2026

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k123906/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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