

K042402 MODIFICATION TO VERTEX RECONSTRUCTION SYSTEMOct 1, 2004
28 days to decisionK042402 · Product code: **KWP** · Orthopedic
Source: <https://www.510kdatabase.net/k042402/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Appliance, Fixation, Spinal Interlaminar (KWP)
Date received	Sep 3, 2004
Decision date	Oct 1, 2004
Days to decision	28 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Medtronic Sofamor Danek, Inc.
Location	Memphis, TN, US
Contact	RICHARD TREHARNE
510(k) history	99 submissions · 89 cleared · 2000-2025

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k042402/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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