

**K090714 VERTEX RECONSTRUCTION SYSTEM, VERTEX  
SELECT RECONSTRUCTION SYSTEM**Apr 17, 2009  
30 days to decisionK090714 · Product code: **KWP** · Orthopedic  
Source: <https://www.510kdatabase.net/k090714/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Appliance, Fixation, Spinal Interlaminar (KWP)
Date received	Mar 18, 2009
Decision date	Apr 17, 2009
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Medtronic Sofamor Danek</b>
Location	Memphis, TN, US
Contact	MELISA LANSKY
510(k) history	154 submissions · 147 cleared · 2002-2021

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k090714/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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