

**K090767 PERPOS FCD-2 SYSTEM (SINGLE PATIENT USE),
ANCHOR, STABILIZER**Jun 11, 2009
80 days to decisionK090767 · Product code: **MRW** · Orthopedic
Source: <https://www.510kdatabase.net/k090767/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent - U
Submission type	Special
Device classification	System, Facet Screw Spinal Device (MRW)
Date received	Mar 23, 2009
Decision date	Jun 11, 2009
Days to decision	80 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Interventional Spine, Inc.
Location	Irvine, CA, US
Contact	CAROL EMERSON
510(k) history	8 submissions · 7 cleared · 2008-2016

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