K090767 PERPOS FCD-2 SYSTEM (SINGLE PATIENT USE), ANCHOR, STABILIZER

CLEARED

Jun 11, 2009
80 days to decision

K090767 · 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