

**K092201 ADVANCE 913 MEDIAL PIVOT TIBIAL INSERT,
ADVANCE 913 MEDIAL PIVOT TIBIAL BASE**Aug 20, 2009
29 days to decisionK092201 · Product code: **JWH** · Orthopedic
Source: <https://www.510kdatabase.net/k092201/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Prosthesis, Knee, Patellofemorotibial, Semi-constrained, Cemented, Polymer/metal/polymer (JWH)
Date received	Jul 22, 2009
Decision date	Aug 20, 2009
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Wrightmedicaltechnologyinc
Location	Arlington, TN, US
Contact	SARAH FAIRFIELD
510(k) history	302 submissions · 291 cleared · 1993-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k092201/>, Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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