

**K092190 SYNTHES DISTRACTION OSTEOGENESIS SYSTEM,
MR CONDITIONAL WITH EXPANDED INDICATIONS**May 11, 2010
294 days to decisionK092190 · Product code: KTT · Orthopedic
Source: <https://www.510kdatabase.net/k092190/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Nail/blade/plate Combination, Multiple Component (KTT)
Date received	Jul 21, 2009
Decision date	May 11, 2010
Days to decision	294 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Synthes (Usa)
Location	Mchenry, IL, US
Contact	JILL R YELTON
510(k) history	411 submissions · 394 cleared · 1977-2015

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k092190/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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