

K051600 SYNTHES (USA) DISTRACTION OSTEOGENESIS SYSTEMJul 27, 2005
41 days to decisionK051600 · Product code: **KTT** · Orthopedic
Source: <https://www.510kdatabase.net/k051600/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Nail/blade/plate Combination, Multiple Component (KTT)
Date received	Jun 16, 2005
Decision date	Jul 27, 2005
Days to decision	41 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k051600/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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