

K033158 SYNTHES REPROCESSED EXTERNAL FIXATION DEVICES

Nov 5, 2003
36 days to decision

K033158 · Product code: **KTT** · Orthopedic
Source: <https://www.510kdatabase.net/k033158/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Nail/blade/plate Combination, Multiple Component (KTT)
Date received	Sep 30, 2003
Decision date	Nov 5, 2003
Days to decision	36 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k033158/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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