

**K950384 SYNTHES(USA) SYNTHES HYBRID EXTERNAL
FIXATOR**May 22, 1995
110 days to decisionK950384 · Product code: LXT · Orthopedic
Source: <https://www.510kdatabase.net/k950384/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Nail/blade/plate Combination, Multiple Component, Metal Composite (LXT)
Date received	Feb 1, 1995
Decision date	May 22, 1995
Days to decision	110 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k950384/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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