

**K073135 SYNTHES (USA) HYBRID EXTERNAL FIXATOR  
MODIFICATIONS**Jan 29, 2008  
83 days to decisionK073135 · Product code: LXT · Orthopedic  
Source: <https://www.510kdatabase.net/k073135/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Nail/blade/plate Combination, Multiple Component, Metal Composite (LXT)
Date received	Nov 7, 2007
Decision date	Jan 29, 2008
Days to decision	83 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Synthes (Usa)</b>
Location	Mchenry, IL, US
Contact	JILL SHERMAN
510(k) history	411 submissions · 394 cleared · 1977-2015

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k073135/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 14, 2026