

K033071 SYNTHES (USA) CANNULATED TITANIUM HUMERAL NAIL SYSTEMNov 5, 2003
37 days to decisionK033071 · Product code: **JDS** · Orthopedic
Source: <https://www.510kdatabase.net/k033071/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Nail, Fixation, Bone (JDS)
Date received	Sep 29, 2003
Decision date	Nov 5, 2003
Days to decision	37 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k033071/> 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