

K042899 SYNTHES (USA) RIA SYSTEM - EXPANDED INDICATIONSMar 16, 2005
147 days to decisionK042899 · Product code: **HTO** · Orthopedic
Source: <https://www.510kdatabase.net/k042899/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Reamer (HTO)
Date received	Oct 20, 2004
Decision date	Mar 16, 2005
Days to decision	147 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k042899/> 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