

K891903 COHN TARGETEERMay 15, 1989
49 days to decisionK891903 · Product code: **LXH** · Orthopedic
Source: <https://www.510kdatabase.net/k891903/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Orthopedic Manual Surgical Instrument (LXH)
Date received	Mar 27, 1989
Decision date	May 15, 1989
Days to decision	49 days
Third-party review	No

APPLICANT

Company	Orthopedic Systems, Inc.
Location	Mchenry, IL, US
Contact	ROBERT R MOORE
510(k) history	95 submissions · 89 cleared · 1977-1996

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k891903/>; 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 21, 2026