FDA 510(k) Premarket Notification

K891903 COHN TARGETEER

CLEARED

May 15, 1989
49 days to decision

K891903 · 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