FDA 510(k) Premarket Notification

K882407 BOW & ARROW DRILL GUIDE

CLEARED

Sep 6, 1988
88 days to decision

K882407 · Product code: **LXI** · Orthopedic
Source: https://www.510kdatabase.net/k882407/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Guide, Drill, Ligament (LXI)
Date received	Jun 10, 1988
Decision date	Sep 6, 1988
Days to decision	88 days
Third-party review	No

APPLICANT

Company	**Instrument Makar, Inc.**
Location	Mchenry, IL, US
Contact	DEAN Z LOOK
510(k) history	28 submissions · 27 cleared · 1979-2000