

**K882407 BOW & ARROW DRILL GUIDE**Sep 6, 1988  
88 days to decisionK882407 · Product code: **LXI** · Orthopedic  
Source: <https://www.510kdatabase.net/k882407/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k882407/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 14, 2026