

**K880739 DYONICS DRILL GUIDE/ISOMETER SYSTEM**Mar 21, 1988  
26 days to decisionK880739 · Product code: **LXI** · Orthopedic  
Source: <https://www.510kdatabase.net/k880739/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Guide, Drill, Ligament (LXI)
Date received	Feb 24, 1988
Decision date	Mar 21, 1988
Days to decision	26 days
Third-party review	No

**APPLICANT**

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Company	<b>Dyonics, Inc.</b>
Location	Walker, MI, US
Contact	KENNETH E CARRIER
510(k) history	19 submissions · 19 cleared · 1978-1990

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k880739/>; 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 22, 2026