

**K950646 INTEGRATOR**Apr 30, 1996  
442 days to decisionK950646 · Product code: **LXM** · Physical Medicine  
Source: <https://www.510kdatabase.net/k950646/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Manipulator, Plunger-like Joint (LXM)
Date received	Feb 13, 1995
Decision date	Apr 30, 1996
Days to decision	442 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Moyco Union Broach, Div., Moyco Technologies, Inc.</b>
Location	York, PA, US
Contact	CHRIS M LEHR
510(k) history	1 submissions · 1 cleared · 1996-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k950646/> 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 June 29, 2026