

**K932236 PRO. HI-LOW TABLE MODEL 9530**Jul 22, 1993  
76 days to decisionK932236 · Product code: **INQ** · Physical Medicine  
Source: <https://www.510kdatabase.net/k932236/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Table, Powered (INQ)
Date received	May 7, 1993
Decision date	Jul 22, 1993
Days to decision	76 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Bailey Mfg. Co.</b>
Location	Lodi, OH, US
Contact	THOMAS CAMPBELL
510(k) history	11 submissions · 11 cleared · 1993-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k932236/> 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