

**K932237 PRO. HI-LOW TREATMENT TABLES MODELS
4050,4070,4090**Jul 22, 1993
76 days to decisionK932237 · Product code: **INQ** · Physical Medicine
Source: <https://www.510kdatabase.net/k932237/>**SUBMISSION DETAILS**

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

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k932237/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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