FDA 510(k) Premarket Notification

K932236 PRO. HI-LOW TABLE MODEL 9530

CLEARED

Jul 22, 1993
76 days to decision

K932236 · Product code: **INQ** · Physical Medicine
Source: https://www.510kdatabase.net/k932236/

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