FDA 510(k) Premarket Notification

K910149 ROMFLEX REHAB EQUIP LATERAL FLEXION UNIT

CLEARED

Jan 18, 1991
4 days to decision

K910149 · Product code: **BXB** · Physical Medicine
Source: https://www.510kdatabase.net/k910149/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Exerciser, Powered (BXB)
Date received	Jan 14, 1991
Decision date	Jan 18, 1991
Days to decision	4 days
Third-party review	No

APPLICANT

Company	**Dyna Flex, Intl.**
Location	Clinton, TN, US
Contact	HARRISON BEAL
510(k) history	4 submissions · 4 cleared · 1990-1991