

K910149 ROMFLEX REHAB EQUIP LATERAL FLEXION UNITJan 18, 1991
4 days to decisionK910149 · Product code: **BXB** · Physical MedicineSource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k910149/>; Data sourced from FDA 510(k) public records (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. - Generated June 30, 2026