FDA 510(k) Premarket Notification

K954508 RETRACTOR

CLEARED

Nov 14, 1995
47 days to decision

K954508 · Product code: **HRX** · Orthopedic
Source: https://www.510kdatabase.net/k954508/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Arthroscope (HRX)
Date received	Sep 28, 1995
Decision date	Nov 14, 1995
Days to decision	47 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Phx Technologies Corp.**
Location	Denton, TX, US
Contact	JAMES F CHAPEL
510(k) history	49 submissions · 49 cleared · 1993-1998

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k954508/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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