FDA 510(k) Premarket Notification

K865066 ARTHROFLATOR

CLEARED

Apr 20, 1987
117 days to decision

K865066 · Product code: **HRX** · Orthopedic
Source: https://www.510kdatabase.net/k865066/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Arthroscope (HRX)
Date received	Dec 24, 1986
Decision date	Apr 20, 1987
Days to decision	117 days
Third-party review	No

APPLICANT

Company	**F.M. Wiest USA, Inc.**
Location	Cherry Hill, NJ, US
Contact	WAYNE W DISANZA
510(k) history	12 submissions · 12 cleared · 1986-1994