FDA 510(k) Premarket Notification

K896770 ARTHRODISTRACTOR

CLEARED

Jan 2, 1990
33 days to decision

K896770 · Product code: **HRX** · Orthopedic
Source: https://www.510kdatabase.net/k896770/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Arthroscope (HRX)
Date received	Nov 30, 1989
Decision date	Jan 2, 1990
Days to decision	33 days
Third-party review	No

APPLICANT

Company	**Arthroplastics, Inc.**
Location	Washington, DC, US
Contact	NORMAN J PHILION
510(k) history	1 submissions · 1 cleared · 1990-1990