

K896770 ARTHRODISTRACTORJan 2, 1990
33 days to decisionK896770 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k896770/>, 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 May 22, 2026