

K900070 MODIFIED USES OF THE ARTHROSCOPIC SURGICAL SYSTEMFeb 16, 1990
43 days to decisionK900070 · Product code: **HRX** · Orthopedic
Source: <https://www.510kdatabase.net/k900070/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Arthroscope (HRX)
Date received	Jan 4, 1990
Decision date	Feb 16, 1990
Days to decision	43 days
Third-party review	No

APPLICANT

Company	Dyonics, Inc.
Location	Walker, MI, US
Contact	ERIC BANNON
510(k) history	19 submissions · 19 cleared · 1978-1990

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k900070/> 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