

K821672 PROFLEXJul 22, 1982
43 days to decisionK821672 · Product code: **HRX** · Orthopedic
Source: <https://www.510kdatabase.net/k821672/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Arthroscope (HRX)
Date received	Jun 9, 1982
Decision date	Jul 22, 1982
Days to decision	43 days
Third-party review	No

APPLICANT

Company	Utah Medical Products, Inc.
Location	Mchenry, IL, US
510(k) history	38 submissions · 38 cleared · 1979-2015

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k821672/> 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 23, 2026