

K830120 IMPROVED ARTHROSCOPEMar 17, 1983
65 days to decisionK830120 · Product code: **HRX** · Orthopedic
Source: <https://www.510kdatabase.net/k830120/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Arthroscope (HRX)
Date received	Jan 11, 1983
Decision date	Mar 17, 1983
Days to decision	65 days
Third-party review	No

APPLICANT

Company	Markoptic
Location	Walker, MI, US
510(k) history	2 submissions · 2 cleared · 1983-1986

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Device record: <https://www.510kdatabase.net/k830120/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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