FDA 510(k) Premarket Notification

K830120 IMPROVED ARTHROSCOPE

CLEARED

Mar 17, 1983
65 days to decision

K830120 · 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