FDA 510(k) Premarket Notification

K821672 PROFLEX

CLEARED

Jul 22, 1982
43 days to decision

K821672 · 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