

K771200 ORTHOSCOPEJul 21, 1977
16 days to decisionK771200 · Product code: **HRX** · Orthopedic
Source: <https://www.510kdatabase.net/k771200/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Arthroscope (HRX)
Date received	Jul 5, 1977
Decision date	Jul 21, 1977
Days to decision	16 days
Third-party review	No

APPLICANT

Company	Stryker Corp.
Location	Mchenry, IL, US
510(k) history	124 submissions · 121 cleared · 1976-2023

Stryker Corp. is an American multinational medical technology company headquartered in Portage, Michigan. The company develops and markets surgical equipment, implants, and patient safety technologies used globally across multiple medical specialties. Stryker has received FDA 510(k) clearances from total submissions since its first clearance in 1976. The company maintains active regulatory engagement, with its latest clearance in 2023. Its product portfolio spans orthopedic devices, neurosurgical implants, surgical instruments, and endoscopy systems, reflecting a broad pr...