FDA 510(k) Premarket Notification

K954465 FMS DUO

CLEARED

Nov 9, 1995
45 days to decision

K954465 · Product code: **HRX** · Orthopedic
Source: https://www.510kdatabase.net/k954465/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Arthroscope (HRX)
Date received	Sep 25, 1995
Decision date	Nov 9, 1995
Days to decision	45 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Future Medical Systems, Inc.**
Location	New York City, NY, US
Contact	PATRICK JANIN
510(k) history	12 submissions · 11 cleared · 1995-2003

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k954465/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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