FDA 510(k) Premarket Notification

K951843 REF. 4102CV INTERMEDIARY TUBING, REF. 4509CV STERILE ZONE KIT

CLEARED

Jul 14, 1995
85 days to decision

K951843 · Product code: **HRX** · Orthopedic
Source: https://www.510kdatabase.net/k951843/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Arthroscope (HRX)
Date received	Apr 20, 1995
Decision date	Jul 14, 1995
Days to decision	85 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