

**K951843 REF. 4102CV INTERMEDIARY TUBING, REF. 4509CV  
STERILE ZONE KIT**Jul 14, 1995  
85 days to decisionK951843 · Product code: **HRX** · Orthopedic  
Source: <https://www.510kdatabase.net/k951843/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k951843/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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