

**K050580 SMITH & NEPHEW INTELIJET FLUID MANAGEMENT SYSTEMS**Apr 20, 2005  
44 days to decisionK050580 · Product code: **HRX** · Orthopedic  
Source: <https://www.510kdatabase.net/k050580/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Arthroscope (HRX)
Date received	Mar 7, 2005
Decision date	Apr 20, 2005
Days to decision	44 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Smith &amp; Nephew Endoscopy, Inc.</b>
Location	Andover, MA, US
Contact	KATHLEEN BURNS
510(k) history	18 submissions · 17 cleared · 1996-2011

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k050580/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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