

**K892376 INTRAVISION(TM) ARTHROSCOPE SYSTEM  
(ADD&apos;L APPLI.)**Aug 1, 1989  
118 days to decisionK892376 · Product code: **HRX** · Orthopedic  
Source: <https://www.510kdatabase.net/k892376/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Arthroscope (HRX)
Date received	Apr 5, 1989
Decision date	Aug 1, 1989
Days to decision	118 days
Third-party review	No

**APPLICANT**

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Company	<b>Concept, Inc.</b>
Location	Mchenry, IL, US
Contact	MARTHA C MILLER
510(k) history	83 submissions · 82 cleared · 1976-1992

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k892376/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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