

**K865066 ARTHROFLATOR**Apr 20, 1987  
117 days to decisionK865066 · Product code: **HRX** · Orthopedic  
Source: <https://www.510kdatabase.net/k865066/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Arthroscope (HRX)
Date received	Dec 24, 1986
Decision date	Apr 20, 1987
Days to decision	117 days
Third-party review	No

**APPLICANT**

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Company	<b>F.M. Wiest USA, Inc.</b>
Location	Cherry Hill, NJ, US
Contact	WAYNE W DISANZA
510(k) history	12 submissions · 12 cleared · 1986-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k865066/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 22, 2026