

**K890695 HI-RIDER WHEELCHAIR**Apr 10, 1989  
59 days to decisionK890695 · Product code: **IPL** · Physical MedicineSource: <https://www.510kdatabase.net/k890695/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wheelchair, Standup (IPL)
Date received	Feb 10, 1989
Decision date	Apr 10, 1989
Days to decision	59 days
Third-party review	No

**APPLICANT**

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Company	<b>Retec USA, Inc.</b>
Location	Orchard Park, NY, US
Contact	FRANK J KOPFER
510(k) history	1 submissions · 1 cleared · 1989-1989

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k890695/>; 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 June 30, 2026