

**K931755 RIWOLITH 2280 PROBES**Apr 15, 1994  
372 days to decisionK931755 · Product code: **FFK** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k931755/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lithotripter, Electro-hydraulic (FFK)
Date received	Apr 8, 1993
Decision date	Apr 15, 1994
Days to decision	372 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Richard Wolf Medical Instruments Corp.</b>
Location	Mchenry, IL, US
Contact	ROBERT L CASARSA
510(k) history	142 submissions · 142 cleared · 1977-2016

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k931755/> 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 14, 2026