

**K860571 F.M. WEIST,KG (URODYNAMIC SYSTEMS)**Mar 18, 1986  
32 days to decisionK860571 · Product code: **FAP** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k860571/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cystometric Gas (carbon-dioxide) On Hydraulic Device (FAP)
Date received	Feb 14, 1986
Decision date	Mar 18, 1986
Days to decision	32 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/k860571/>, 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