

**K880342 UROSTHESIA**Feb 25, 1988  
30 days to decisionK880342 · Product code: **FMF** · General HospitalSource: <https://www.510kdatabase.net/k880342/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Jan 26, 1988
Decision date	Feb 25, 1988
Days to decision	30 days
Third-party review	No

**APPLICANT**

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Company	<b>Medical Research Innovators, Inc.</b>
Location	El Cajon, CA, US
Contact	SEAN ROWLAND
510(k) history	1 submissions · 1 cleared · 1988-1988

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k880342/>; 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 July 9, 2026