

**K122785 HYDROSIL**May 14, 2013  
245 days to decisionK122785 · Product code: **EZD** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k122785/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Straight (EZD)
Date received	Sep 11, 2012
Decision date	May 14, 2013
Days to decision	245 days
Third-party review	No
Summary / Statement	Summary
Other names	MAGIC; PERSONAL CATHETER

**APPLICANT**

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Company	<b>Rochester Medical Corp.</b>
Location	Rochester, MN, US
Contact	ROB ANGLIN
510(k) history	13 submissions · 13 cleared · 1990-2013

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k122785/>; 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 9, 2026