

**K042224 RM3 RENAL PRESERVATION SYSTEM**Oct 22, 2004  
66 days to decisionK042224 · Product code: **KDN** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k042224/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, Perfusion, Kidney (KDN)
Date received	Aug 17, 2004
Decision date	Oct 22, 2004
Days to decision	66 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Waters Instruments, Inc.</b>
Location	Walker, MI, US
Contact	DAVE SCHOLLMAN
510(k) history	12 submissions · 12 cleared · 1981-2006

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k042224/> 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 26, 2026