

**K130192 NEXTSTEP RETROGRADE**May 10, 2013  
105 days to decisionK130192 · Product code: **MSD** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k130192/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - K
Submission type	Traditional
Device classification	Catheter, Hemodialysis, Implanted (MSD)
Date received	Jan 25, 2013
Decision date	May 10, 2013
Days to decision	105 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Arrow International, Inc.</b>
Location	Reading, PA, US
Contact	GWEN TASCHNER
510(k) history	19 submissions · 17 cleared · 2003-2020

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k130192/> 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 18, 2026