

**K123630 FRESENIUS LIBERTY CYCLER**Sep 9, 2013  
290 days to decisionK123630 · Product code: **FKX** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k123630/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Peritoneal, Automatic Delivery (FKX)
Date received	Nov 23, 2012
Decision date	Sep 9, 2013
Days to decision	290 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Fresenius Medical Care North America, Design Cente</b>
Location	Waltham,, MA, US
Contact	DENISE OPPERMANN
510(k) history	5 submissions · 5 cleared · 2010-2013

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k123630/> 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 8, 2026