

**K091360 CLEARFLUX DIALYZER REPROCESSING SYSTEM**Nov 3, 2010  
544 days to decisionK091360 · Product code: **LIF** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k091360/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dialyzer Reprocessing System (LIF)
Date received	May 8, 2009
Decision date	Nov 3, 2010
Days to decision	544 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Novaflex Technologies</b>
Location	Princeton, NJ, US
Contact	MOHAMED E LABIB
510(k) history	1 submissions · 1 cleared · 2010-2010

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k091360/> 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 20, 2026