

**K112142 DIMENSION VISTA LITHIUM FLEX REAGENT  
CARTRIDGE**Nov 4, 2011  
101 days to decisionK112142 · Product code: **NDW** · Chemistry  
Source: <https://www.510kdatabase.net/k112142/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Assay, Porphyrin, Spectrophotometry, Lithium (NDW)
Date received	Jul 26, 2011
Decision date	Nov 4, 2011
Days to decision	101 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Siemens Healthcare Diagnostics</b>
Location	Newark, DE, US
Contact	ROSE T MARINELLI
510(k) history	92 submissions · 92 cleared · 2008-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k112142/> 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 28, 2026