

**K882232 DISCOVER(R)**Oct 25, 1988  
151 days to decisionK882232 · Product code: **LCX** · Chemistry  
Source: <https://www.510kdatabase.net/k882232/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Kit, Test, Pregnancy, Hcg, Over The Counter (LCX)
Date received	May 27, 1988
Decision date	Oct 25, 1988
Days to decision	151 days
Third-party review	No

**APPLICANT**

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Company	<b>Carter Products</b>
Location	Cranbury, NJ, US
Contact	STEPHEN KOLAKOWSKY
510(k) history	20 submissions · 20 cleared · 1988-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k882232/>; 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 29, 2026