

K171136 LIA Pregnancy TestNov 20, 2017
217 days to decisionK171136 · Product code: **LCX** · Chemistry
Source: <https://www.510kdatabase.net/k171136/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Kit, Test, Pregnancy, Hcg, Over The Counter (LCX)
Date received	Apr 17, 2017
Decision date	Nov 20, 2017
Days to decision	217 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Lia Diagnostics
Location	Philadelphia, PA, US
Contact	Anna Couturier
510(k) history	1 submissions · 1 cleared · 2017-2017

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k171136/> Data sourced from FDA 510(k) public records (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. - Generated May 31, 2026