FDA 510(k) Premarket Notification

# K171136 LIA Pregnancy Test

**CLEARED**

Nov 20, 2017
217 days to decision

K171136 · 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 |