

**K993141 AT HOME QUICKCUP PREGNANCY TEST, MODEL 9015**Nov 8, 1999  
48 days to decisionK993141 · Product code: **LCX** · Chemistry  
Source: <https://www.510kdatabase.net/k993141/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Kit, Test, Pregnancy, Hcg, Over The Counter (LCX)
Date received	Sep 21, 1999
Decision date	Nov 8, 1999
Days to decision	48 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Phamatech</b>
Location	San Diego, CA, US
Contact	CARL A MONGIOVI
510(k) history	47 submissions · 47 cleared · 1996-2004

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k993141/> 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 June 28, 2026