FDA 510(k) Premarket Notification

K993141 AT HOME QUICKCUP PREGNANCY TEST, MODEL 9015

CLEARED

Nov 8, 1999
48 days to decision

K993141 · 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	**Phamatech**
Location	San Diego, CA, US
Contact	CARL A MONGIOVI
510(k) history	47 submissions · 47 cleared · 1996-2004