

**K982621 QUICKSCREEN AT HOME DRUG TEST, MODEL
9149/9150**Oct 16, 1998
80 days to decisionK982621 · Product code: **MVO** · Toxicology
Source: <https://www.510kdatabase.net/k982621/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Kit, Test, Multiple, Drugs Of Abuse, Over The Counter (MVO)
Date received	Jul 28, 1998
Decision date	Oct 16, 1998
Days to decision	80 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k982621/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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