FDA 510(k) Premarket Notification

K992685 QUICKSCREEN WORKPLACE DRUG SCREENING TEST, MODEL 9147 25C

CLEARED

Oct 6, 1999
57 days to decision

K992685 · Product code: **MVO** · Toxicology
Source: https://www.510kdatabase.net/k992685/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Kit, Test, Multiple, Drugs Of Abuse, Over The Counter (MVO)
Date received	Aug 10, 1999
Decision date	Oct 6, 1999
Days to decision	57 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.net
Device record: https://www.510kdatabase.net/k992685/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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