FDA 510(k) Premarket Notification

K993165 QUICKSCREEN PRO MULTI DRUG SCREEN, MODEL 9177 & 9178

CLEARED

Oct 20, 1999
28 days to decision

K993165 · Product code: **DJG** · Toxicology
Source: https://www.510kdatabase.net/k993165/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Enzyme Immunoassay, Opiates (DJG)
Date received	Sep 22, 1999
Decision date	Oct 20, 1999
Days to decision	28 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/k993165/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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