FDA 510(k) Premarket Notification

K971218 QUIKSTRIP ONE STEP AMPHETAMINE TEST

CLEARED

May 30, 1997
58 days to decision

K971218 · Product code: **DKZ** · Toxicology
Source: https://www.510kdatabase.net/k971218/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Enzyme Immunoassay, Amphetamine (DKZ)
Date received	Apr 2, 1997
Decision date	May 30, 1997
Days to decision	58 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Drial Consultants, Inc.**
Location	Menlo Park, CA, US
Contact	CLEVE W LAIRD
510(k) history	13 submissions · 13 cleared · 1997-1998

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k971218/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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