FDA 510(k) Premarket Notification

K972059 QUIKPAC II ONE STEP COCAINE TEST

CLEARED

Aug 28, 1997
87 days to decision

K972059 · Product code: **DIO** · Toxicology
Source: https://www.510kdatabase.net/k972059/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Enzyme Immunoassay, Cocaine And Cocaine Metabolites (DIO)
Date received	Jun 2, 1997
Decision date	Aug 28, 1997
Days to decision	87 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/k972059/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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