FDA 510(k) Premarket Notification

K000979 REDI-SCREEN

CLEARED

Apr 10, 2000
14 days to decision

K000979 · Product code: **LAF** · Toxicology
Source: https://www.510kdatabase.net/k000979/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Gas Chromatography, Methamphetamine (LAF)
Date received	Mar 27, 2000
Decision date	Apr 10, 2000
Days to decision	14 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Redwood Biotech, Inc.**
Location	Sunnyvale, CA, US
Contact	JANIS FREESTONE
510(k) history	6 submissions · 6 cleared · 1999-2000

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k000979/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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