

**K013002 TRIAGE TOX DRUG SCREEN CALIBRATION
VERIFICATION CONTROLS, CATALOG #94002**Oct 3, 2001
27 days to decisionK013002 · Product code: **DIF** · Toxicology
Source: <https://www.510kdatabase.net/k013002/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drug Mixture Control Materials (DIF)
Date received	Sep 6, 2001
Decision date	Oct 3, 2001
Days to decision	27 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Biosite Incorporated
Location	San Diego, CA, US
Contact	JEFFREY R DAHLEN
510(k) history	46 submissions · 45 cleared · 1990-2010

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k013002/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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