

**K012999 TRIAGE TOX DRUG SCREEN CONTROLS, CATALOG  
#94001**Oct 3, 2001  
27 days to decisionK012999 · Product code: **DIF** · Toxicology  
Source: <https://www.510kdatabase.net/k012999/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k012999/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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