

**K954322 TRI-POINT LIQUIMMUNE LIQUID ASSAYED
IMMUNOASSAY CONTROL**Oct 11, 1995
26 days to decisionK954322 · Product code: **DIF** · Toxicology
Source: <https://www.510kdatabase.net/k954322/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drug Mixture Control Materials (DIF)
Date received	Sep 15, 1995
Decision date	Oct 11, 1995
Days to decision	26 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Medical Analysis Systems, Inc.
Location	Mchenry, IL, US
Contact	KEVIN KENNAN
510(k) history	107 submissions · 107 cleared · 1978-2004

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k954322/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 14, 2026