FDA 510(k) Premarket Notification

K954322 TRI-POINT LIQUIMMUNE LIQUID ASSAYED IMMUNOASSAY CONTROL

CLEARED

Oct 11, 1995
26 days to decision

K954322 · 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