

K932753 ROCHE ABUSCREEN TRI-LEVEL REFERENCE CONTROLSJul 22, 1993
55 days to decisionK932753 · Product code: **DKB** · Toxicology
Source: <https://www.510kdatabase.net/k932753/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Calibrators, Drug Mixture (DKB)
Date received	May 28, 1993
Decision date	Jul 22, 1993
Days to decision	55 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Roche Diagnostic Systems, Inc.
Location	Mchenry, IL, US
Contact	RITA SMITH
510(k) history	296 submissions · 296 cleared · 1983-1999

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k932753/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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