

**K052053 LIQUICHEK QUALITATIVE URINE TOXICOLOGY
CONTROL, NEGATIVE, POSITIVE, MINIPAK**Aug 9, 2005
11 days to decisionK052053 · Product code: **DIF** · Toxicology
Source: <https://www.510kdatabase.net/k052053/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drug Mixture Control Materials (DIF)
Date received	Jul 29, 2005
Decision date	Aug 9, 2005
Days to decision	11 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Bio-Rad
Location	Mchenry, IL, US
Contact	SUZANNE S PARSONS
Website	http://www.bio-rad.com
510(k) history	319 submissions · 319 cleared · 1976-2017

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k052053/> 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