

**K971691 LIQUICHEK URINE TOXICOLOGY CONTROL - LEVEL  
S1, S2, S3**May 30, 1997  
23 days to decisionK971691 · Product code: **DIF** · Toxicology  
Source: <https://www.510kdatabase.net/k971691/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drug Mixture Control Materials (DIF)
Date received	May 7, 1997
Decision date	May 30, 1997
Days to decision	23 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Bio-Rad</b>
Location	Mchenry, IL, US
Contact	ELIZABETH PLATT
Website	<a href="http://www.bio-rad.com">http://www.bio-rad.com</a>
510(k) history	319 submissions · 319 cleared · 1976-2017

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k971691/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://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](https://accessdata.fda.gov). - Generated May 14, 2026