

**K914170 LIQUICHEK THERAPEUTIC DRUG MONITORING CONTROL(TDM)**Nov 22, 1991  
66 days to decisionK914170 · Product code: **DIF** · Toxicology  
Source: <https://www.510kdatabase.net/k914170/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drug Mixture Control Materials (DIF)
Date received	Sep 17, 1991
Decision date	Nov 22, 1991
Days to decision	66 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>Bio-Rad</b>
Location	Mchenry, IL, US
Contact	SHARON K.SMTH,PH.D
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/k914170/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 14, 2026