

K042836 LIQUICHEK RETICULOCYTE CONTROL (A)Nov 5, 2004
22 days to decisionK042836 · Product code: **JPK** · Hematology
Source: <https://www.510kdatabase.net/k042836/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mixture, Hematology Quality Control (JPK)
Date received	Oct 14, 2004
Decision date	Nov 5, 2004
Days to decision	22 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Bio-Rad
Location	Mchenry, IL, US
Contact	ELIZABETH PLATT
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/k042836/> 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