FDA 510(k) Premarket Notification

K042836 LIQUICHEK RETICULOCYTE CONTROL (A)

CLEARED

Nov 5, 2004
22 days to decision

K042836 · 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
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510(k) history	319 submissions · 319 cleared · 1976-2017